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(12) **United States Patent**
Desarbre et al.(10) **Patent No.:** **US 9,346,799 B2**(45) **Date of Patent:** ***May 24, 2016**(54) **USEFUL COMBINATIONS OF
MONOBACTAM ANTIBIOTICS WITH
BETA-LACTAMASE INHIBITORS**(71) Applicant: **Basilea Pharmaceutica AG**, Basel (CH)(72) Inventors: **Eric Desarbre**, Mulhouse (FR);
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(CH)(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.This patent is subject to a terminal dis-
claimer.(21) Appl. No.: **14/512,847**(22) Filed: **Oct. 13, 2014**(65) **Prior Publication Data**

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2006, now Pat. No. 8,901,293.(30) **Foreign Application Priority Data**Dec. 7, 2005 (EP) 05026699
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A61K 31/427 (2006.01)
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C07D 401/14 (2006.01)
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(2013.01); **A61K 31/41** (2013.01); **A61K**
31/427 (2013.01); **A61K 31/43** (2013.01);
A61K 31/433 (2013.01); **A61K 31/4375**
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C07D 487/04 (2013.01)(58) **Field of Classification Search**USPC 514/210.05
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See application file for complete search history.(56) **References Cited**

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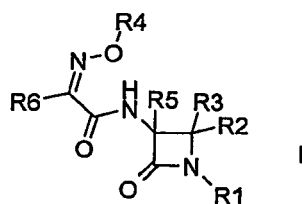
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(57) **ABSTRACT**A pharmaceutical composition, comprising a combination of
an antibiologically active compound of the formula (I): with a
 β -lactamase inhibitor of one of the formulae (II) to (XIII) are
active against Gram-negative bacteria, in particular such bac-
teria which have become resistant against antibiotics such as
aztreonam, carumonam and tigemonam. Optionally the com-
positions may comprise another β -lactamase inhibitor of one
of the formulae (II) to (XIII), particularly of formula (V) or
formula (VI).



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USEFUL COMBINATIONS OF MONOBACTAM ANTIBIOTICS WITH BETA-LACTAMASE INHIBITORS

This application is a divisional application of U.S. application Ser. No. 12/086,270, filed Sep. 22, 2009, which, in turn, is a National Stage Application of PCT/CH2006/000685 filed Dec. 7, 2006, which claims priority from European Patent Application 06006291.6, filed Mar. 27, 2006, and European Patent Application 05026699.8 filed Dec. 7, 2005, the contents of all of which are expressly incorporated herein by reference. Priority of the PCT and European Patent Applications is claimed.

FIELD OF THE INVENTION

The present invention is concerned with pharmaceutical compositions and methods for treating infections caused by pathogenic Gram-negative bacteria.

BACKGROUND OF THE INVENTION

β -Lactam antibiotics have been widely used for the treatment of bacterial infections both in hospitals and in the general public. There are several classes of β -lactam antibiotics that have found clinical application, these include the penicillins, cephalosporins, cephamycins, carbacephems, oxacephems, carbapenems and monobactams.

The efficiency of all of these classes to cure bacterial infections has been impaired by the appearance of bacteria that are resistant towards the antibiotics. The prevalent cause of this resistance in Gram-negative bacteria is the expression by the bacteria of enzymes known as β -lactamases that are able to hydrolyse the β -lactam antibiotics rendering them inactive. Bacteria are able to produce a variety of β -lactamases, including penicillinases, cephalosporinases, cephamycinases, carbapenemases, monobactamases, broad-spectrum β -lactamases and extended-spectrum β -lactamases.

The possibility of rescuing individual β -lactam antibiotics by combination with a β -lactamase inhibitor that inactivates the β -lactamase before it can hydrolyse the β -lactam antibiotic has been demonstrated with clinically useful combinations between penicillins such as amoxicillin, ampicillin and ticarcillin and β -lactamase inhibitors such as clavulanic acid, sulbactam and tazobactam. Further, potential combinations have been described involving cephalosporins and newly developed β -lactamase inhibitors including bridged monobactams, penam sulfones, phosphonate esters, exomethylene penams and diazabicyclooctane derivatives.

Monobactams have been regarded as stable towards many β -lactamases. However, there are now many strains of Gram-negative bacteria that exhibit β -lactamase-mediated resistance towards the monobactam antibiotics (aztreonam, carumonam and tigemonam).

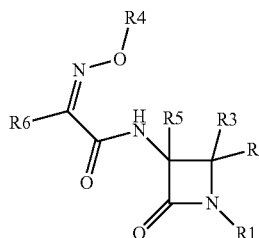
The present invention aims to provide improved medications with novel monobactam antibiotics and combinations of monobactam antibiotics with β -lactamase inhibitors that are active against aerobic Gram-negative bacteria that are resistant against treatments with monobactam antibiotics.

SUMMARY OF THE INVENTION

The objective set is solved by a pharmaceutical composition, comprising a combination of

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a) an antibiotically active compound of the following formula I:



in which

R1 signifies SO_3H , OSO_3H , $\text{CRaRa}'\text{COOH}$, $\text{OCRaRa}'\text{COOH}$, 5-tetrazolyl, SO_2NHRb or CONHRc ,

wherein Ra and Ra' are independently selected from hydrogen; alkyl; allyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkoxyalkyl and a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

wherein Rb is hydrogen; alkyl; alkoxycarbonyl; alkylaminocarbonyl; benzylaminocarbonyl in which the benzyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or phenylaminocarbonyl in which the phenyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

wherein Rc is hydrogen; alkyl; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkoxy-carbonyl; SO_2phenyl ; $\text{SO}_2\text{NHalkyl}$; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

R2 and R3 independently signify hydrogen; alkyl; alkenyl; alkynyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; azido; halogen; dihalogenomethyl; trihalogenomethyl; alkoxycarbonyl; carboxyl; sulfonyl or $\text{CH}_2\text{X1}$,

wherein X1 is azido; amino; halogen; hydroxyl; cyano; carboxyl; aminosulfonyl; alkoxycarbonyl; alkanoylamino; phenylaminocarbonyl; alkylaminocarbonyl; aminocarbonyl; carbamoyloxy; alkylaminosulfonyl; phenylaminosulfonyl in which the phenyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or benzyl which may be substituted with 1 to 5 substituents

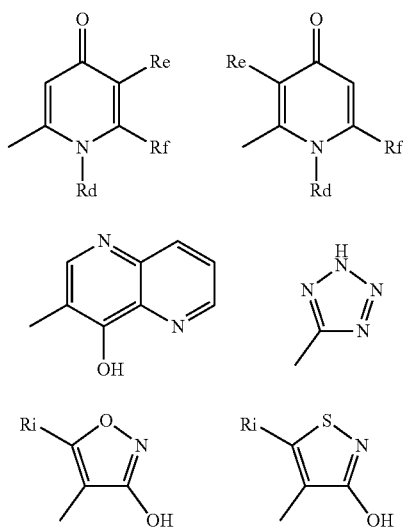
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selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;
R4 signifies hydrogen; alkyl; C(Rx)(Ry)Z,

wherein Rx and Ry are independently selected from hydrogen; alkyl; allyl; (C₃-C₆)cycloalkyl; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; (C₂-C₇) alkene and (C₂-C₇)alkyne; or Rx and Ry taken together may form an alkylene bridge —(CH₂)_n— with n being an integer number from 2 to 6; and

Z is COOH; CH₂N(OH)COR' wherein

R' is hydrogen, alkyl, alkylamino, alkoxy, benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen, phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen, or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or Z is one of the following six groups



in which groups

Rd, Re and Rf are independently selected from hydrogen; alkyl; amino; monoalkylamino; carboxylaminoalkyl; alkoxycarbonyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; diphenylmethyl; trityl; and ORg wherein

Rg is hydrogen; alkyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino and halogen; or phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino and halogen;

or, when Re and Rf are vicinal substituents, Re and Rf taken together may also be —O—CH=CH—CH₂—, —O—CH₂—CH₂—O—, —CH₂—CH₂—CH₂—, —CH₂—CH₂—CH₂—CH₂—, —CH=CH—CH=CH— or CH=C(OH)—C(OH)=CH—;

Ri is hydrogen; alkyl; alkylamino; alkoxy; benzyl which may be substituted with 1 to 5 substituents selected

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from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl and hydroxyl; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

R5 signifies hydrogen, alkyl, halogenomethyl, dihalogenomethyl, trihalogenomethyl, alkoxy, formylamino or alkylcarbonylamino;

R6 signifies phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, carbonylamino and halogen;

or a pharmaceutically acceptable salt thereof;

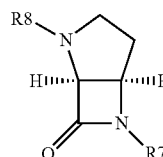
and one or more β-lactamase inhibitors selected from the following groups b1) to b11):

b1) a bridged monobactam derivative of the following formula II:

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II

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in which:

R7 signifies SO₃H, OSO₃H or OCRjRj'COOH,

wherein Rj and Rj' are independently selected from hydrogen; alkyl; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkylamino and alkoxyalkyl;

R8 is alkoxycarbonylamino, the acyl residue of an α or β-amino acid, or a residue of the formula Q-(X)_r-Y—,

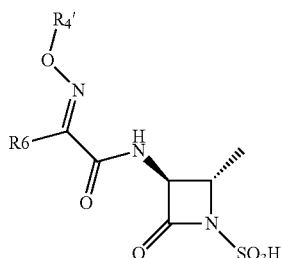
wherein Q is a 3-6 membered ring which optionally contains nitrogen, sulphur and/or oxygen and which is optionally fused to a phenyl ring or to a 5-6 membered heterocyclic ring and which is optionally substituted with 1 to 4 substituents selected from alkyl, allyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino, carboxamide which may be substituted, carboxylic acid, carbonylalkoxy, aminocarbonyl, alkylaminocarbonyl, halogen, halogenomethyl, dihalogenomethyl, trihalogenomethyl, sulfamide, substituted sulfamide with substituents selected from alkyl, phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino and halogen and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, halogen and benzyl, urea which may be substituted with alkyl, aminoalkyl or alkoxy and carbamate which may be substituted with alkyl, aminoalkyl or alkoxy,

X signifies a linear spacer of from 1 to 6 atoms length and containing carbon, nitrogen, oxygen and/or sulphur atoms, of which up to 2 atoms can be nitrogen atoms and 1 atom can be oxygen or sulphur,

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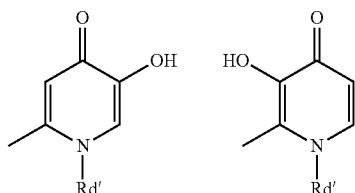
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r is an integer of from 0 to 1; and
Y is selected from —CO—, —CS—, —NHCO—, —NH—
CONH— and —SO₂—;
or a pharmaceutically acceptable salt thereof;
or
b2) a monobactam derivative of the general formula III:

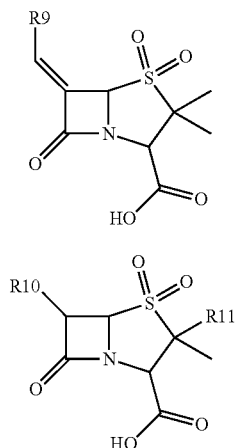


in which

R4' signifies hydrogen, alkyl or CH(Rx')Z', wherein
Rx' is selected from hydrogen; (C₁-C₆)alkyl; allyl; phenyl
and (C₃-C₆)cycloalkyl; and Z' signifies COOH or a
group of one of the following two formulae:



in which Rd' is hydrogen or hydroxy; and
R6 is as defined for formula I; or a pharmaceutically accept-
able salt thereof;
or
b3) a penam sulfone derivative of the general formulae IV or
V:



in which

R9 signifies COOH or a 5-6 membered monocyclic or
polycyclic heteroaromatic group;
R10 signifies hydrogen or halogen;
R11 signifies CH₂R12; CH=CHR12 wherein R12 is
hydrogen, halogen, cyano, carboxylic acid, acyl such as
acetyl, carboxamide which may be substituted, alkoxy-
carbonyl or a 5-6 membered heteroaromatic ring which
is optionally substituted with 1 to 5 substituents selected
from alkyl, hydroxyl, alkoxy, amino, alkylamino,
dialkylamino and halogen; or which is optionally fused

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with a 5-6 membered heteroaromatic ring; CH=NR12'
wherein R12' is amino, alkylamino, dialkylamino, ami-
nocarbonyl, acylamino such as acetylamino, hydroxy,
alkoxy,

5 or a pharmaceutically acceptable salt thereof;
or
b4) an oxapenam derivative of the general formula VI:

III

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in which

R13 signifies OR14; S(O)_nR14 or a 5-6 membered het-
eroaromatic ring which may be substituted with 1 to 5
substituents selected from alkyl, hydroxyl, alkoxy,
amino, alkylamino, dialkylamino and halogen; whereby
n=0, 1 or 2, and R14 is hydrogen, alkyl, (C₂-C₇)alkene,
(C₂-C₇)alkyne or a 5-6 membered heteroaromatic ring
which may be substituted with 1 to 5 substituents
selected from alkyl, hydroxyl, alkoxy, amino, alky-
lamino, dialkylamino and halogen,

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or a pharmaceutically acceptable salt thereof;

or

b5) a penem derivative of the general formula VII:

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IV

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V

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in which

R15 signifies a 5-6 membered heteroaromatic ring which
may be substituted with 1 to 5 substituents selected from
alkyl, hydroxyl, alkoxy, amino, alkylamino, dialky-
lamino and halogen; or which is optionally fused with a
5-6 membered heteroaromatic ring and/or which is
optionally bound to the exo-methylene group over a
—CH=CH— spacer being preferably in the (E)-con-
figuration,

or a pharmaceutically acceptable salt thereof;

or

b6) a cephem sulfone derivative of the general formula VIII:

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in which

R16 signifies COOR17, whereby R17 signifies hydrogen
or alkyl; or a 5-6 membered heteroaromatic ring which is
optionally fused with a 5-6 membered heteroaromatic
ring being optionally substituted with 1 to 5 substituents

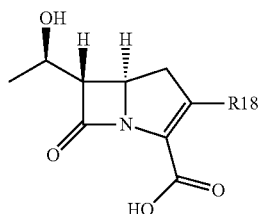
VI

VII

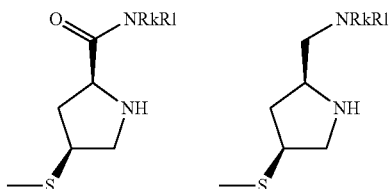
VIII

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selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino, halogen; and/or being optionally bound to the exo-methylene group over a —CH=CH— spacer being preferably in the (E)-configuration, or a pharmaceutically acceptable salt thereof;
or
b7) a carbapenem derivative of the general formula IX:



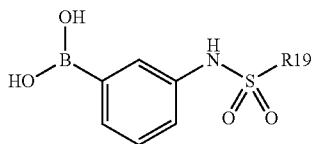
in which R18 signifies —S-alkyl , $\text{—S—(CH}_2\text{)}_2\text{—NH—CH=NH}$ or a group of the following two formulae



wherein Rk and Rl are individually selected from hydrogen, alkyl, 2-, 3-, 4-carboxyphenyl and sulfamoyl, or a pharmaceutically acceptable salt thereof;

or

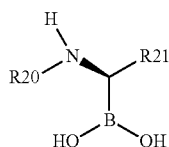
b8) a boronate derivative of the general formula X:



wherein R19 signifies a 5-6 membered heteroaromatic ring which may be substituted with amino, alkylamino, dialkylamino or alkylsulfoxide, or a pharmaceutically acceptable salt thereof;

or

b9) a boronate derivative of the general formula XI:

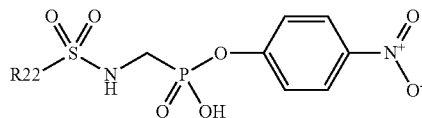


wherein

R20 and 21 are independently selected from a 5-6 membered heteroaromatic ring or phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and

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halogen and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen, or a pharmaceutically acceptable salt thereof;
5 or
b10) a phosphonate derivative of the general formula XII:



XII

IX 10

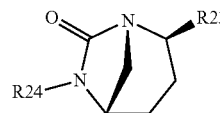
15 wherein

R22 is selected from a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen and which is optionally fused with a 5-6 membered heteroaromatic ring; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen,

or a pharmaceutically acceptable salt thereof;

or

b11) a diazabicyclooctane derivative of the general formula XIII:



XIII

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X 40

in which

R23 signifies hydrogen, carboxylic acid, alkoxycarbonyl or carboxamide which may be substituted, and

R24 signifies SO_3H , OSO_3H or OCRjRj'COOH , wherein Rj and Rj' are as defined for formula II,

or a pharmaceutically acceptable salt thereof.

As a preferred embodiment of the invention, the pharmaceutical compositions may comprise two or more compounds selected from one of the formulae II to XIII of b1) to b11), these two or more compounds being different from each other.

The object set is also solved by the novel monobactam antibiotics of formula Ia as described hereinafter, which may be used in the same combinations as outlined above.

Further objects of the invention are visible from the description hereinafter and from the appended claims.

DETAILED DESCRIPTION OF THE INVENTION

It has surprisingly been found that the efficacy of monobactam antibiotics of the formula I against aerobic Gram-negative bacteria can be potentiated by co-using a β -lactamase inhibitor according to any one of the formulae II to XIII.

In formula I, when the oxazetidine ring is lying in the plane of the paper, preferably R3 points downwards from the plane, and R2 points upwards from the plane.

For the purposes of the present invention, the compounds of the above formula III are not considered as “antibiotically active” compounds in the sense of claim 1. The compounds in formula I have the oximino group predominantly the “syn” configuration shown in formula I, whereas the compounds III have the oximino group specifically in the “anti” configuration shown in formula III.

The term “alkyl”, as used in the present application, preferably means straight-chain or branched (C₁-C₇)alkyl such as in particular methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, isopropyl, isobutyl, tert-butyl or neopentyl.

The term “alkoxy”, as used in the present application, preferably means straight-chain or branched (C₁-C₇)alkoxy, such as in particular methoxy, ethoxy, propoxy, 1- or 2-butoxy, 1-, 2-, or 3-pentyloxy, 1-, 2- or 3-hexyloxy, 1-, 2-, 3-, or 4-heptyloxy or tert-butoxy. The term “alkylhydroxyl” as also used, shall be considered synonymous to “alkoxy”; in particular for the alkyl in “alkylhydroxyl” the same definition shall apply as given above for “alkyl”.

The term “carboxamide which may be substituted” preferably has the meaning that the carboxamide has 0 to 2 hydrogen atoms attached to the amino moiety, with the remainder of the substituents on the amino moiety being alkyl or phenyl which may be substituted.

The term “imine which may be substituted” preferably means that the imine bears at the imine nitrogen hydrogen, alkyl, phenyl which may be substituted or benzyl which may be substituted.

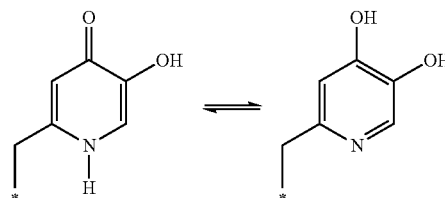
The terms “optionally substituted phenyl” and “optionally substituted benzyl”, if given without specifically indicated substituents, shall preferably mean that the phenyl or benzyl is optionally substituted with 1 to 5 substituents selected from alkyl, alkoxy, dialkylamino and halogen, wherein the “alkyl” itself and the alkyl in dialkylamino and alkoxy has the meaning as defined above.

The term “linear spacer”, as used in the present application, preferably means a linear divalent group selected from —O—, —S—, —NH—, —NH—NH—, —CH₂—, —CO—, —CH₂O—, —CH₂CH₂—, —CH=CH—, —CH₂NH—, —S—CH₂—, —SO₂—, —CH₂—, —O—CH₂—, —S—CH₂CH₂—, —CH₂CH₂—NH—, —CH₂—NH—CO—CH₂CH₂—, —CH₂—NH—CO—O—

CH₂CH₂, —CH₂—NH—CO—NH—CH₂CH₂, —CH₂—O—CO—NH—CH₂CH₂—, —CH(OH)—, —CH(COOH)—, —CH(OSO₃H)—, —CH(OCONH₂)— and —CH[CH(CH₃)₂]—.

Some compounds of formula I may, when they contain an acidic group (such as when R1 is SO₃H, OSO₃H, CRaRa'COOH, OCRaRa'COOH or SO₂NHRb) be present as a salt with a pharmaceutically acceptable inorganic base (e.g. NaOH, KOH, NH₃, K₂CO₃, Na₂CO₃, Na₂HPO₄ or K₂HPO₄) or organic base (e.g. NEt₃, HNiPr₂, triethanolamine, TRIS or basic amino acids such as arginine and lysine). The use of such salts of compounds of formula I is encompassed by the invention. Also, some of the compounds of formula I, when they contain both an acidic group (such as when R1 is SO₃H, OSO₃H, CRaRa'COOH, OCRaRa'COOH or SO₂NHRb) and a basic group (such as when R6 is 2-amino-1,3-thiazol-4-yl, 5-amino-1,2,4-thiadiazol-3-yl, 5-amino-1,2,4-oxadiazol-3-yl, 3-aminoisoxazol-5-yl, 5-amino-1-methylpyrazol-3-yl, 5-aminopyrazol-3-yl, 6-amino-2-pyridyl, 4-aminopyrimidin-2-yl, 2-carbonylamino-1,3-thiazol-4-yl or 2-amino-5-chloro-1,3-thiazol-4-yl) may form an inner, zwitterionic salt; such inner salts are also intended to be encompassed by the claims.

When R4 contains a 4-pyridone moiety, wherein Rd is selected from hydrogen and hydroxy, there is the possibility of tautomerism:



The invention intends to encompass the use of any such tautomers.

A first group of preferred examples of compounds of the formula I for the combinations of the invention are aztreonam, carumonam, tigemonam, and compounds according to the following table 1 (R5 is in these compounds always H):

TABLE 1

Compound number	R1	R2	R3	R4	R6
1	SO ₃ H	H	CH ₃		
2	SO ₃ H	H	CH ₃		

TABLE 1-continued

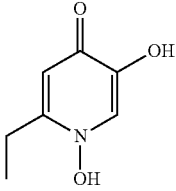
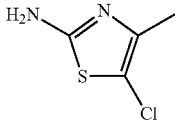
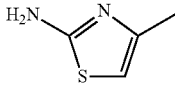
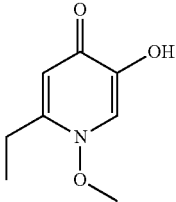
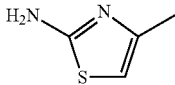
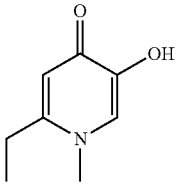
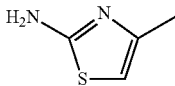
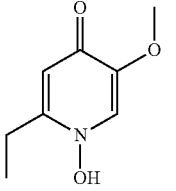
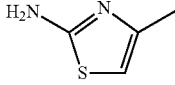
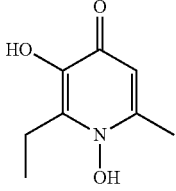
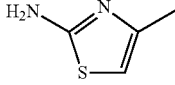
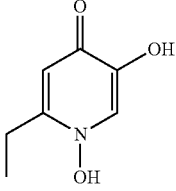
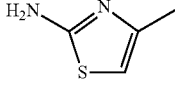
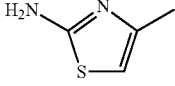
Compound number	R1	R2	R3	R4	R6
3	SO ₃ H	H	CH ₃		
4	SO ₃ H	H	CH ₃	CH ₃	
<u>5</u>	SO ₃ H	H	CH ₃		
<u>6</u>	SO ₃ H	H	CH ₃		
<u>7</u>	SO ₃ H	H	CH ₃		
<u>8</u>	SO ₃ H	H	CH ₃		
<u>9</u>	SO ₃ H	H	H		
<u>10</u>	SO ₃ H	H	CH ₃	CH ₂ CH ₂ N(OH)COCH ₃	

TABLE 1-continued

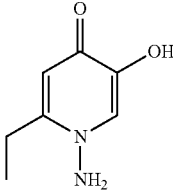
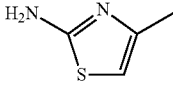
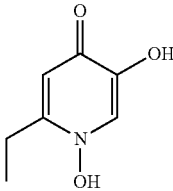
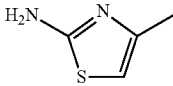
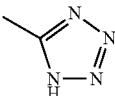
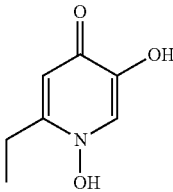
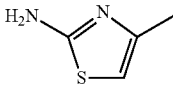
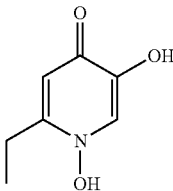
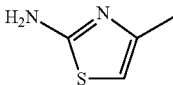
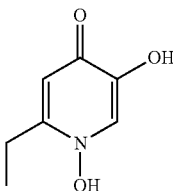
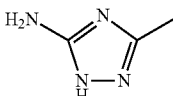
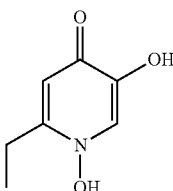
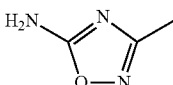
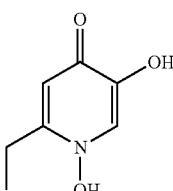
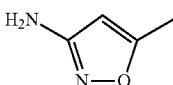
Compound number	R1	R2	R3	R4	R6
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<u>12</u>	OSO ₃ H	H	CH ₃		
<u>13</u>		H	CH ₃		
<u>14</u>	SO ₂ NH ₂	H	CH ₃		
<u>15</u>	SO ₃ H	H	CH ₃		
<u>16</u>	SO ₃ H	H	CH ₃		
<u>17</u>	SO ₃ H	H	CH ₃		

TABLE 1-continued

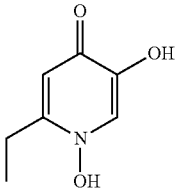
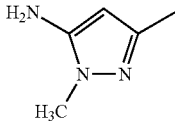
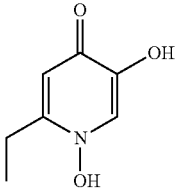
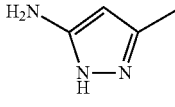
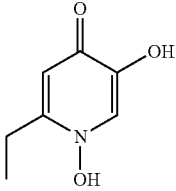
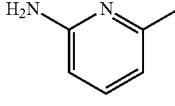
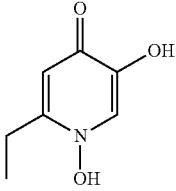
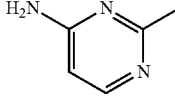
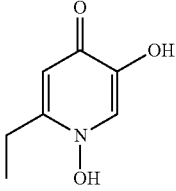
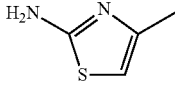
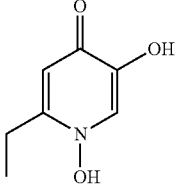
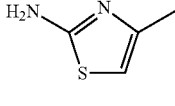
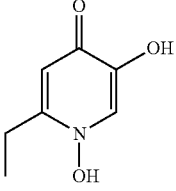
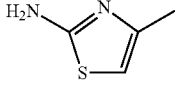
Compound number	R1	R2	R3	R4	R6
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<u>19</u>	SO ₃ H	H	CH ₃		
<u>20</u>	SO ₃ H	H	CH ₃		
<u>21</u>	SO ₃ H	H	CH ₃		
<u>22</u>	OCH ₂ COOH	H	CH ₃		
<u>23</u>	OCH(CH ₃)COOH	H	CH ₃		
<u>24</u>	OSO ₃ H	CH ₂ F	H		

TABLE 1-continued

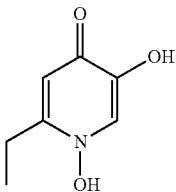
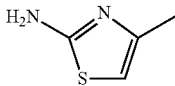
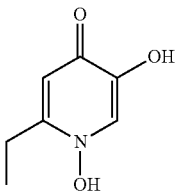
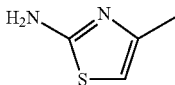
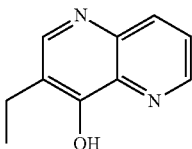
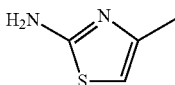
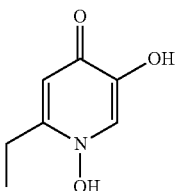
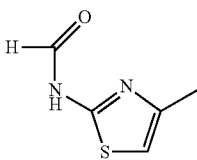
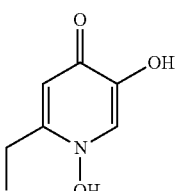
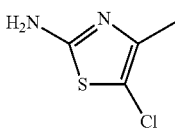
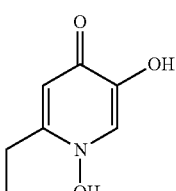
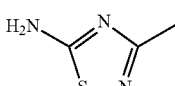
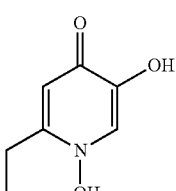
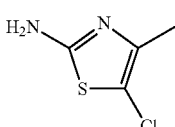
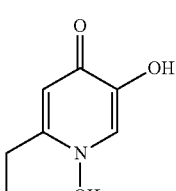
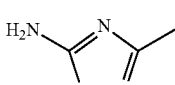
Compound number	R1	R2	R3	R4	R6
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<u>26</u>	OSO ₃ H	CH ₃	CH ₃		
<u>27</u>	SO ₃ H	H	CH ₃		
<u>28</u>	SO ₃ H	H	CH ₃		
<u>29</u>	OSO ₃ H	H	CH ₃		
<u>30</u>	OSO ₃ H	H	CH ₃		
<u>31</u>	OSO ₃ H	CH ₃	CH ₃		
<u>32</u>	OSO ₃ H	CH ₃	CH ₃		

TABLE 1-continued

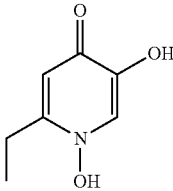
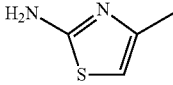
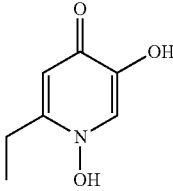
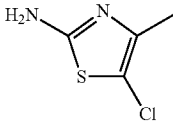
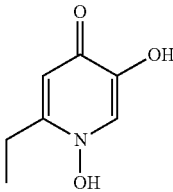
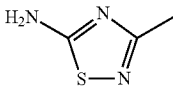
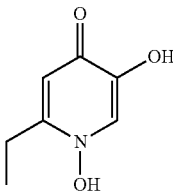
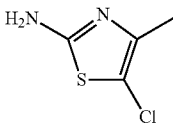
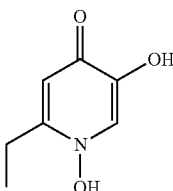
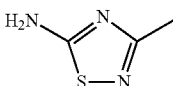
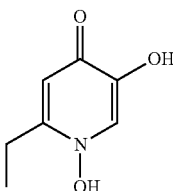
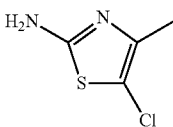
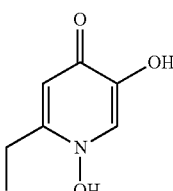
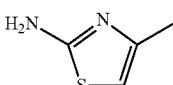
Compound number	R1	R2	R3	R4	R6
<u>33</u>	OCH ₂ COOH	CH ₃	CH ₃		
<u>34</u>	OCH ₂ COOH	CH ₃	CH ₃		
<u>35</u>	OCH ₂ COOH	CH ₃	CH ₃		
<u>36</u>	OCH ₂ COOH	H	CH ₃		
<u>37</u>	OCH ₂ COOH	H	CH ₃		
<u>38</u>	SO ₃ H	CH ₃	CH ₃		
<u>39</u>	SO ₃ H	CH ₃	CH ₃		

TABLE 1-continued

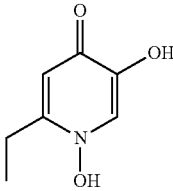
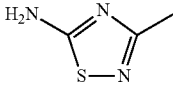
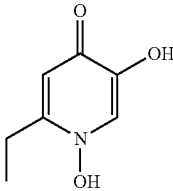
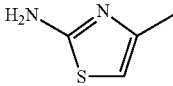
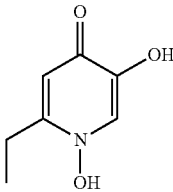
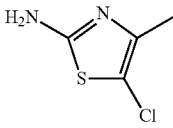
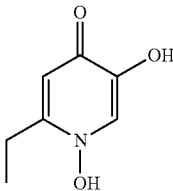
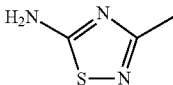
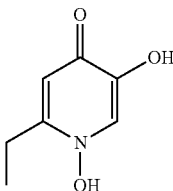
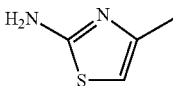
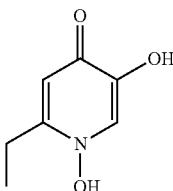
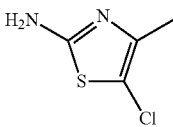
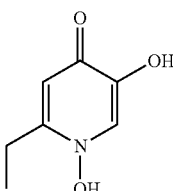
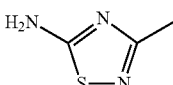
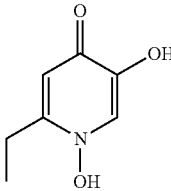
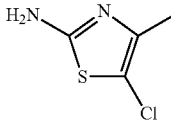
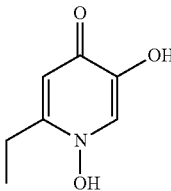
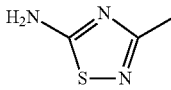
Compound number	R1	R2	R3	R4	R6
<u>40</u>	SO ₃ H	CH ₃	CH ₃		
<u>41</u>	OCH ₂ COOH	CH ₂ OCONH ₂	H		
<u>42</u>	OCH ₂ COOH	CH ₂ OCONH ₂	H		
<u>43</u>	OCH ₂ COOH	CH ₂ OCONH ₂	H		
<u>44</u>	OCH ₂ COOH	CH ₂ F	H		
<u>45</u>	OCH ₂ COOH	CH ₂ F	H		
<u>46</u>	OCH ₂ COOH	CH ₂ F	H		

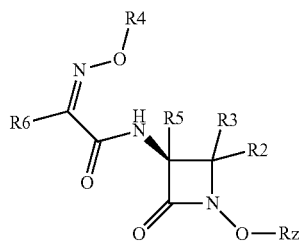
TABLE 1-continued

Compound number	R1	R2	R3	R4	R6
<u>47</u>	OSO ₃ H	CH ₂ OCONH ₂	H		
<u>48</u>	OSO ₃ H	CH ₂ OCONH ₂	H		

The numbering of the compounds as given in above table 1 is used in the following for the sake of conciseness.

The compounds of the above table 1 are, when their compound numbers are underscored and bold, part of the present invention.

Among the compounds of formula I which are more preferred in the combinations of the invention and which are novel per se are those of the following formula Ia:

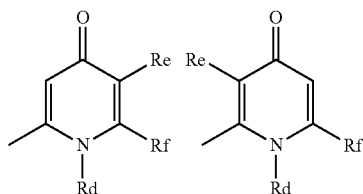


wherein

Rz is SO₃H or CRaRa'COOH, wherein Ra and Ra' are as defined for formula I;

R2, R3, R5 and R6 are as defined for formula I;

R4 is CH₂Z; whereby Z is a group of one of the formulae



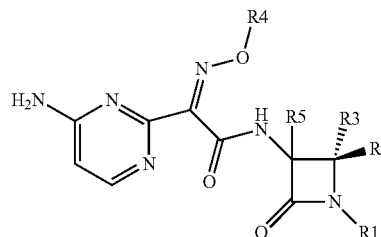
wherein Rd, Re and Rf are as defined for the compounds of formula I; and the pharmaceutically acceptable salts or inner salts thereof. This first group of compounds forms part of the invention. When Rz is SO₃H, then preferably both R2 and R3 are methyl. When Rz is CRaRa'COOH, then R2 is preferably selected from hydrogen, methyl, fluoromethyl and carbamoyloxymethyl; and R3 is preferably selected from or hydrogen and methyl; and more preferably here, the absolute configuration at the carbon atom bearing R2 and R3 is (S).

Preferably, Ra and Ra' are each hydrogen. Preferably for all compounds of formula Ia, the Rd, Re and Rf are individually selected from hydrogen and hydroxy, with the proviso that at least two of Rd, Re and Rf are hydroxy (most preferably Rd and Re are hydroxy and Rf is hydrogen). R5 is preferably hydrogen. R6 is preferably an optionally amino-substituted and optionally chloro-substituted 5-6-membered heteroaromatic ring, this ring being more preferably selected from 2-amino-1,3-thiazol-4-yl, 5-amino-1,2,4-thiadiazol-3-yl, 5-amino-1,2,4-oxadiazol-3-yl, 3-aminoisoxazol-5-yl, 5-amino-1-methylpyrazol-3-yl, 5-aminopyrazol-3-yl, 6-amino-2-pyridyl, 4-aminopyrimidin-2-yl, 2-carbamoylamino-1,3-thiazol-4-yl, 2-amino-5-chloro-1,3-thiazol-4-yl and 2-thienyl.

More preferred examples of the compound of formula Ia are the compounds (12), (22), (23), (24), (25), (26), (29), (30), (31), (32), (33), (34), (35), (36), (37), (41), (42), (43), (44), (45), (46), (47) and (48) of above table 1. The most preferred compounds of formula Ia are compounds (22), (23), (26) and (31).

The compounds of formula Ia, if Rz is SO₃H, can be made by a methodology as outlined in scheme 4 below, until the R2,R3-disubstituted 3-amino-2-oxoazetidine hydroxysulfonate, and reacting this further in a manner known per se to connect the 3-amino substituents. The compounds of formula Ia, if Rz is CRaRa'COOH, can be prepared following the synthesis scheme of compounds II to X as described in U.S. Pat. No. 4,939,253 (column 15, line 26 to column 17, line 25), and reacting the obtained oxoazetidine X further according to scheme 1 described hereinafter.

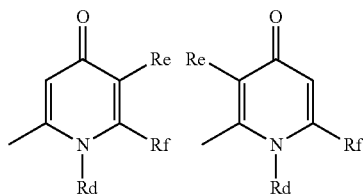
A second group of compounds of formula I which is preferred in the combinations of the invention and is novel are those of the following formula Ib:



and the pharmaceutically acceptable salts thereof, wherein the residues R1, R2, R3, R4 and R5 are as defined for formula

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I. This second group of compounds also forms part of the invention. Preferably, in these compounds R1 is SO₃H, R2 is H, R3 is methyl and R4 is C(Rx)(Ry)Z; whereby Rx=Ry=H, and Z is a group of one of the formulae



wherein R_d, R_e and R_f are individually selected from hydrogen and hydroxy, with the proviso that at least two of R_d, R_e and R_f are hydroxy (most preferably R_d and R_e are hydroxy and R_f is hydrogen).

The antibiotically active monobactams of formula I may firstly be combined with

b1) β -Lactamase Inhibitors of the Above General Formula II.

In formula II, R8 is preferably of the formula Q-(X)_r-CO—. The following subgroups are more preferable within this formula:

a) With X=—CH₂— and r=1; wherein Q is a pyridinium group which may be substituted with one to three substituents, preferably one to two substituents, selected from alkyl; perfluoroalkyl, in particular trifluoromethyl; phenyl; benzyl; R_uR_vN—, wherein R_u and R_v are independently selected from hydrogen, alkyl, cycloalkyl, pyrrolidinyl, carbamoyl and N-(carbamoylalkyl)carbamoyl, or wherein R_u and R_v taken together form an alkylene bridge —(CH₂)_w—, with w being an integer number of 3 to 6; alkylcarbonyl; R_uR_vNCO—, wherein R_u and R_v are as defined before; (alkoxycarbonyl) alkyl; thiocarbamoyl and alkoxycarbonyl; or wherein Q is a pyridinium group which is fused with a 5-6 membered carbocycle; and thioamide. Examples within this subgroup are pyridinium, 2-, 3- or 4-aminopyridinium, 3-N-methylaminopyridinium, 3-N,N-dimethylaminopyridinium, 4-(N-methylamino)pyridinium, 4-(N,N-dimethylamino)pyridinium, 3-carbamoylpyridinium, 3-(N-methylcarbamoyl)pyridinium, 3-(N,N-dimethylcarbamoyl)pyridinium, 4-carbamoylpyridinium, 4-(N-methylcarbamoyl)pyridinium, 4-(N,N-dimethylcarbamoyl)pyridinium, 3-(N-cyclopropylcarbamoyl)pyridinium, 4-(N-cyclopropylcarbamoyl)pyridinium, 4-(N-methylcarbamoyl)pyridinium, 3-(methoxycarbonyl)pyridinium, 3-(ethoxycarbonyl)pyridinium, 4-(methoxycarbonyl)pyridinium, 4-(ethoxycarbonyl)pyridinium, 3-thiocarbamoylpyridinium, 3-(N-methylthiocarbamoyl)pyridinium, 3-(N,N-dimethylthiocarbamoyl)pyridinium, 4-thiocarbamoylpyridinium, 4-(N-methylthiocarbamoyl)pyridinium, 4-(N,N-dimethylthiocarbamoyl)pyridinium, 2,3-, 2,4-, 2,5-, 2,6-, 3,4-, 3,5-, 3,6-dimethylpyridinium, 3- or 4-isopropylpyridinium, 3- or 4-(trifluoromethyl)pyridinium, 3- or 4-phenylpyridinium, 3- or 4-benzylpyridinium, quinolinium, isoquinolinium, 5,6,7,8-tetrahydroquinolinium and 5,6,7,8-tetrahydroisoquinolinium. In this subgroup, R7 is preferably —SO₂[−] or —OSO₂[−], to form a pharmaceutically acceptable inner salt. The compounds II of this subgroup a) themselves are per se also part of the invention, but compound 102 of table 2 below is known from J. Med. Chem. 1998, 41(21), 3961, and does per se not form part of the invention.

b) With X=—SCH₂— and r=1; wherein Q is 1,2,3,4-tetra-

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Examples for this substituent are methyl, ethyl, propyl, butyl, 2-aminoethyl, 2-(N-methylamino)ethyl, 2-(N,N-dimethylamino)ethyl, 3-aminopropyl, 3-(N-methylamino)propyl, 3-(N,N-dimethylamino)propyl, 2-(N-ethylamino)ethyl, 2-(N,N-diethylamino)ethyl, 3-(N-ethylamino)propyl and 3-(N,N-diethylamino)propyl. In these compounds, R7 is preferably —SO₃H or OSO₃H, whereby the possibility of formation of an inner, acid addition salt may be allowed, to form a pharmaceutically acceptable inner salt.

c) With X=—CH₂NH₂— and r=1; wherein Q is phenyl which may be substituted with one to two substituents selected from hydroxy and alkoxy, or a 5-6-membered heterocycle which preferably is selected from oxazol-2-yl, -3-yl or -4-yl, furan-2-yl or 3-yl, thiophen-2-yl or -3-yl, 1,3-thiazol-2-yl, -3-yl, -4-yl or -5-yl and which optionally may be substituted with one to two substituents selected from alkyl and alkoxy.

d) With X=—NH₂— and r=1, wherein Q is phenyl which may be substituted with one to two substituents selected from hydroxy; alkoxy and a substituted urea of the formula H₂N[(CH₂)_mO]_n(CH₂)_oHNCONH— or a substituted carbamate of the formula H₂N[(CH₂)_mO]_n(CH₂)_oHNCOO—, wherein m and o are independently integer numbers from 2 to 3 and n is an integer number from 0 to 1. Examples for Q here are phenyl, 3- and 4-hydroxyphenyl, 2,3-, 2,4-, 2,6-, 3,4-, 3,5- and 3,6-dihydroxyphenyl, 2-, 3- and 4-methoxyphenyl, 3-[N'-(2-(2-aminoethoxy)ethyl)carbamoylamino]phenyl, 3-[N'-(2-(3-aminopropoxy)ethyl)carbamoylamino]phenyl, 3-[N'-(3-(2-aminoethoxy)propyl)carbamoylamino]phenyl, 3-[N'-(3-(3-aminopropoxy)propyl)carbamoylamino]phenyl, 4-[N'-(2-(2-aminoethoxy)ethyl)carbamoylamino]phenyl, 4-[N'-(2-(3-aminopropoxy)ethyl)carbamoylamino]phenyl, 4-[N'-(3-(2-aminoethoxy)propyl)carbamoylamino]phenyl, 4-[N'-(3-(3-aminopropoxy)propyl)carbamoylamino]phenyl, 3-[N'-(2-(2-aminoethoxy)ethyl)carbamoyloxy]phenyl, 3-[N'-(2-(3-aminopropoxy)ethyl)carbamoyloxy]phenyl, 3-[N'-(3-(2-aminoethoxy)propyl)carbamoyloxy]phenyl, 3-[N'-(3-(3-aminopropoxy)propyl)carbamoyloxy]phenyl, 4-[N'-(2-(2-aminoethoxy)ethyl)carbamoyloxy]phenyl, 4-[N'-(2-(3-aminopropoxy)ethyl)carbamoyloxy]phenyl, 4-[N'-(3-(2-aminoethoxy)propyl)carbamoyloxy]phenyl and 4-[N'-(3-(3-aminopropoxy)propyl)carbamoyloxy]phenyl.

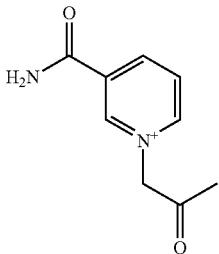
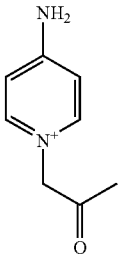
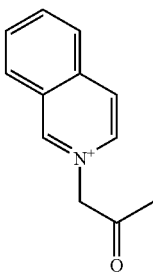
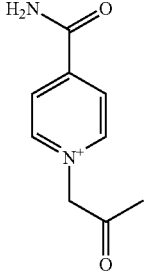
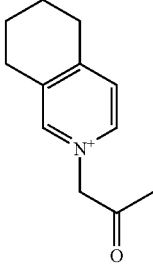
Particularly preferred examples of the compounds of formula II are according to the following table 2

TABLE 2

Compound number	R7	R8
101	SO ₃ [−]	

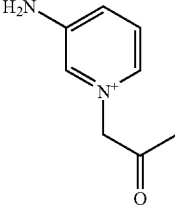
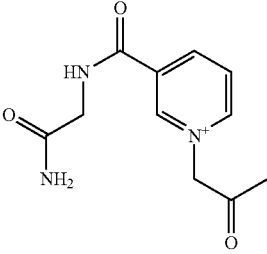
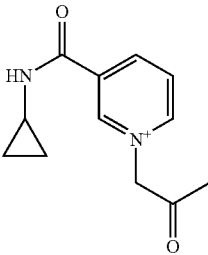
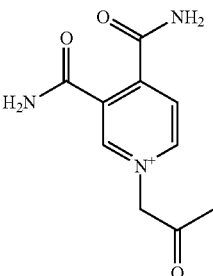
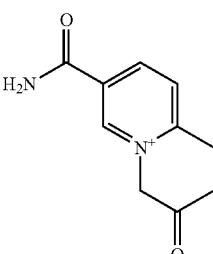
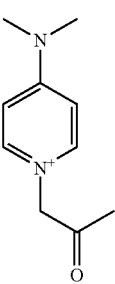
27

TABLE 2-continued

Compound number	R7	R8
102	SO ₃ ⁻	
<u>103</u>	SO ₃ ⁻	
<u>104</u>	SO ₃ ⁻	
<u>105</u>	SO ₃ ⁻	
<u>106</u>	SO ₃ ⁻	

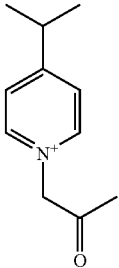
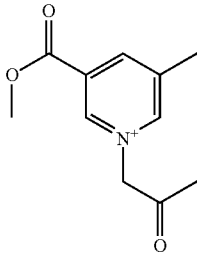
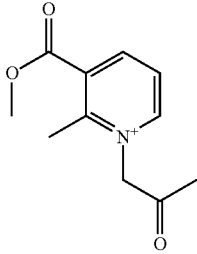
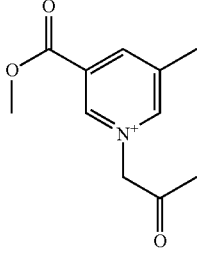
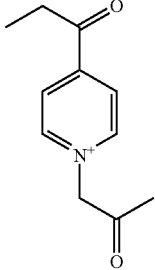
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TABLE 2-continued

Compound number	R7	R8
5 <u>107</u>	SO ₃ ⁻	
10		
15 <u>108</u>	SO ₃ ⁻	
20		
25 <u>109</u>	SO ₃ ⁻	
30		
35 <u>110</u>	SO ₃ ⁻	
40		
45		
50 <u>111</u>	SO ₃ ⁻	
55		
60 <u>112</u>	SO ₃ ⁻	
65		

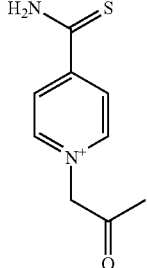
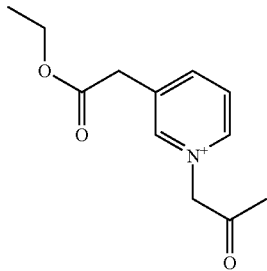
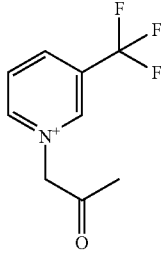
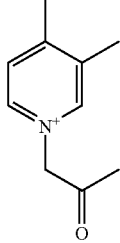
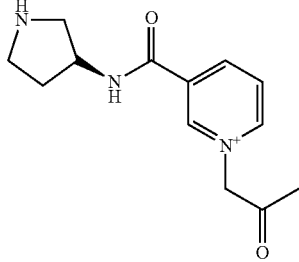
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TABLE 2-continued

Compound number	R7	R8
<u>113</u>	SO ₃ ⁻	
<u>114</u>	SO ₃ ⁻	
<u>115</u>	SO ₃ ⁻	
<u>116</u>	SO ₃ ⁻	
<u>117</u>	SO ₃ ⁻	

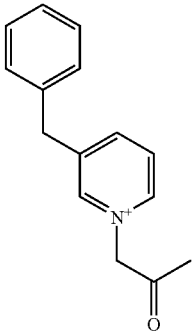
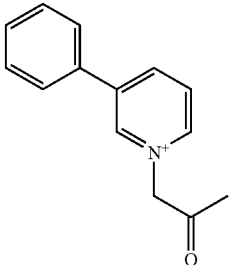
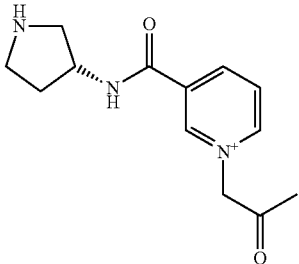
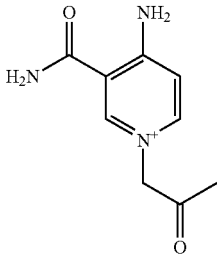
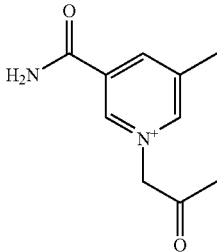
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TABLE 2-continued

Compound number	R7	R8
<u>118</u>	SO ₃ ⁻	
<u>119</u>	SO ₃ ⁻	
<u>120</u>	SO ₃ ⁻	
<u>121</u>	SO ₃ ⁻	
<u>122</u>	SO ₃ ⁻	

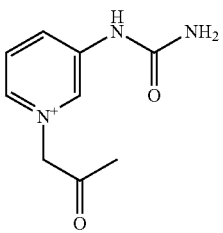
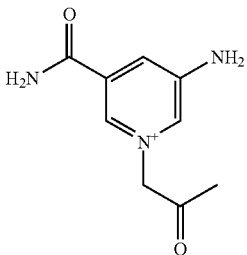
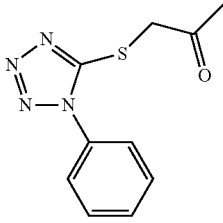
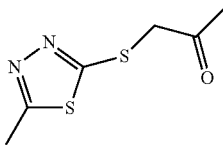
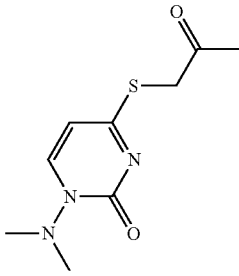
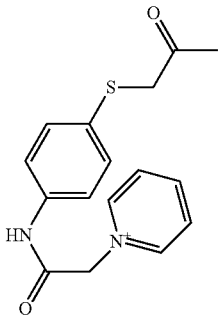
31

TABLE 2-continued

Compound number	R7	R8
<u>123</u>	SO ₃ ⁻	
<u>124</u>	SO ₃ ⁻	
<u>125</u>	SO ₃ ⁻	
<u>126</u>	SO ₃ ⁻	
<u>127</u>	SO ₃ ⁻	

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TABLE 2-continued

Compound number	R7	R8
<u>128</u>	SO ₃ ⁻	
<u>129</u>	SO ₃ ⁻	
201	SO ₃ Na	
202	SO ₃ Na	
203	SO ₃ H	
204	SO ₃ ⁻	

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TABLE 2-continued

Compound number	R7	R8
205	SO ₃ H	
206	SO ₃ Na	
<u>207</u>	SO ₃ H	
<u>208</u>	SO ₃ H	
<u>209</u>	SO ₃ H	
<u>210</u>	SO ₃ H	
<u>211</u>	SO ₃ H	

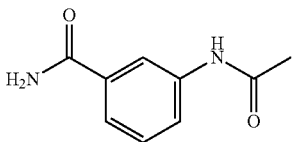
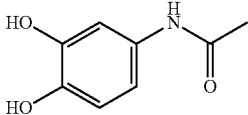
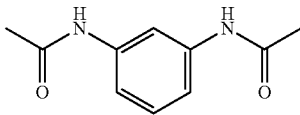
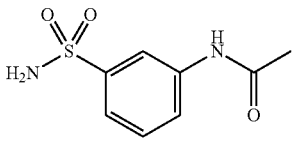
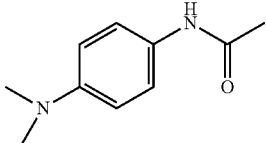
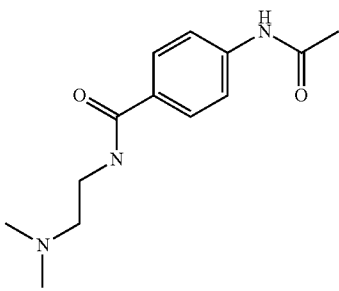
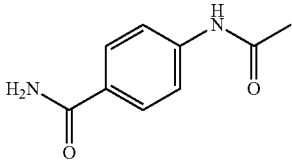
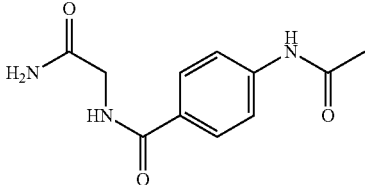
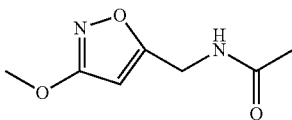
34

TABLE 2-continued

Compound number	R7	R8
<u>212</u>	SO ₃ H	
10		
15	<u>213</u>	SO ₃ H
20		
<u>214</u>	SO ₃ H	
25		
30	<u>215</u>	SO ₃ H
35		
40		
301	SO ₃ Na	
45		
302	SO ₃ Na	
50		
303	SO ₃ Na	
55		
60		
304	SO ₃ Na	
65		

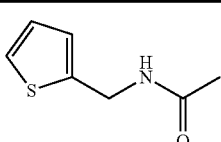
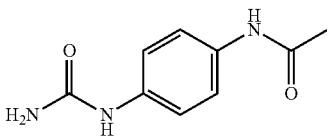
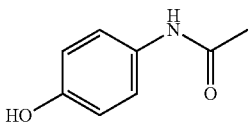
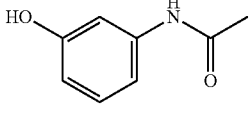
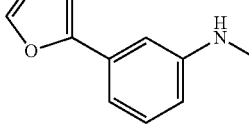
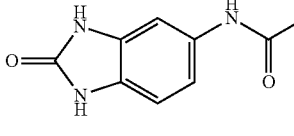
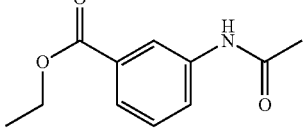
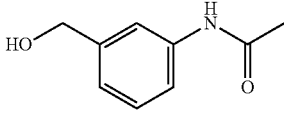
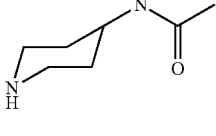
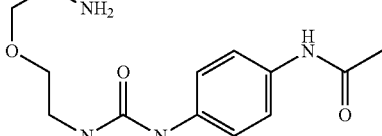

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TABLE 2-continued

Compound number	R7	R8
305	SO ₃ Na	
306	SO ₃ Na	
307	SO ₃ Na	
308	SO ₃ Na	
309	SO ₃ Na	
310	SO ₃ Na	
311	SO ₃ Na	
312	SO ₃ Na	
313	SO ₃ Na	

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TABLE 2-continued

Compound number	R7	R8
5	SO ₃ Na	
10	SO ₃ Na	
15	SO ₃ Na	
20	SO ₃ Na	
25	SO ₃ Na	
30	SO ₃ H	
35	SO ₃ Na	
40	SO ₃ Na	
45	SO ₃ Na	
50	SO ₃ Na	
55	SO ₃ H	
60	SO ₃ H	
65	SO ₃ H	

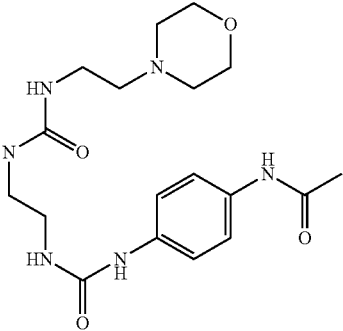
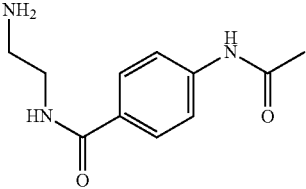
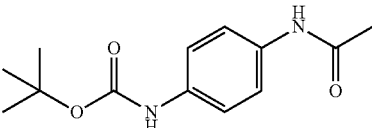
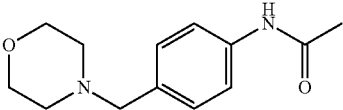
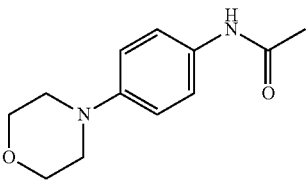
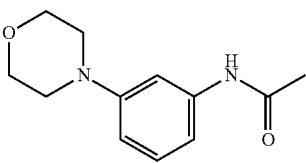
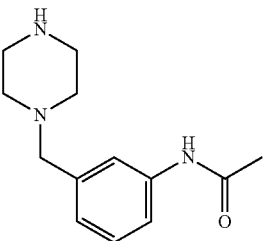
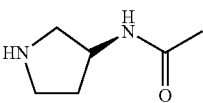
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TABLE 2-continued

Compound number	R7	R8
324	SO ₃ H	
325	SO ₃ H	
326	SO ₃ H	
327	SO ₃ Na	
328	SO ₃ Na	
329	SO ₃ Na	
330	SO ₃ H	

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TABLE 2-continued

Compound number	R7	R8
331	SO ₃ H	
332	SO ₃ H	
333	SO ₃ Na	
<u>334</u>	SO ₃ H	
<u>335</u>	SO ₃ H	
<u>336</u>	SO ₃ H	
<u>337</u>	SO ₃ H	
<u>338</u>	SO ₃ H	

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TABLE 2-continued

Compound number	R7	R8
<u>339</u>	SO ₃ H	
<u>340</u>	SO ₃ H	
<u>341</u>	SO ₃ H	
<u>342</u>	SO ₃ H	
<u>343</u>	SO ₃ H	
<u>344</u>	SO ₃ H	
<u>401</u>	SO ₃ H	
<u>402</u>	SO ₃ H	

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TABLE 2-continued

Compound number	R7	R8
<u>403</u>	SO ₃ H	
<u>404</u>	SO ₃ Na	
<u>405</u>	SO ₃ Na	
<u>406</u>	SO ₃ H	
<u>407</u>	SO ₃ ⁻	
<u>408</u>	SO ₃ Na	

The numbering of the preferred compounds of formula II as given in above table 2 is used in the following for the sake of conciseness.

The compounds of the above table 2 are, when their compound numbers are underscored and bold, part of the present invention. Otherwise they are disclosed in EP-A-0-508 234, U.S. Pat. No. 6,566,355 and J. Med. Chem. 1998, 3961. These publications are hereby incorporated by reference.

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The compounds of formula I may also be combined with other β -lactamase inhibitors. These further inhibitors are:
b2) Monobactam Derivatives of the Above General Formula III:

Examples of these compounds are disclosed in WO-A-99/10324 and WO-A-98/47895, incorporated herein by reference.

A preferred example of these compounds is 3-[(2E)-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl)methoxy]-2-(2-thienyl)-3-azaprop-2-enoylamino](3S,4S)-4-methyl-2-oxoazetidinesulfonic acid disclosed in WO-A-98/47895.

b3) Penam Sulfone Derivatives of the Above General Formulae IV and V:

Here, preferred examples of the 5-6 membered heteroaromatic ring as R12 are 1,3-thiazol-2-yl, 1,2,4-oxadiazol-3-yl and 1,2,3-triazol-1-yl.

Particularly preferred examples of the compounds of formula V are sulbactam, tazobactam and the compounds of the following table (in parentheses the source):

R10	R11	Compound number
hydrogen	(1Z)-2-cyanovinyl	501 EP-A-0 640 607
hydrogen	(1E)-3-oxo-but-1-en-1-yl	502 EP-A-0 640 607
hydrogen	(1Z)-2-(1,3-thiazol-2-yl)vinyl	503 EP-A-0 640 607
hydrogen	(1E)-2-(1,2,4-oxadiazol-3-yl)vinyl	504 EP-A-0 640 607
carboxymethylene	CH ₃	(Bioorg. Med. Chem. Lett. 1995, 1513)
hydrogen	(1E)-2-methoxy-2-azavinyl	US-A-5,686,441
hydrogen	(1E)-2,3-diaza-4-oxo-pent-1-en-1-yl	US-A-5,686,441

The numbers used in the rightmost column of this table are also used in the tests for biological activity (see below).

b4) Oxapenam Derivatives of the Above General Formula VI:

A preferred compound of formula VI is clavulanic acid or a customary pharmaceutically acceptable salt thereof (i.e. a clavulanate).

b5) Penem Derivatives of the Above General Formula VII:

Here, preferred examples of the 5-6 membered heteroaromatic ring which may be bound over a $-\text{CH}=\text{CH}-$ spacer as R15 are 1,2,3-triazol-4-yl, 2H,3H-imidazo[2,1-b]1,3-thiazolidin-6-yl and 2'-[1-methyl-1,2,3-triazolin-4-yl]vinylidene.

Preferred examples for inhibitors of formula VII are e.g. 6-[(1-methyl(1,2,3-triazol-4-yl))methylene]-5-oxo-6aH-azetidino[2,1-b]1,3-thiazoline-3-carboxylic acid and 6-(2H,3H-imidazo[2,1-b]1,3-thiazolidin-6-ylmethylene)-5-oxo-6aH-azetidino[2,1-b]1,3-thiazoline-3-carboxylic acid (as described in Antimicrob. Agents Chemother. 1989, 1580 and Antimicrob. Agents Chemother. 1991, 1748), and 6-[(2E)-3-(1-methyl(1,2,3-triazolin-4-yl))prop-2-enylidene]-5-oxo-6aH-azetidino[2,1-b]1,3-thiazoline-3-carboxylic acid (as described in J. Antibiotic 1997, 50, 350).

b6) Cephem Sulfone Derivatives of the Above General Formula VIII:

Here, a preferred example of the 5-6 membered heteroaromatic ring as R16 is 2-thienyl.

Preferred examples are here 3-(acetyloxymethyl)-1,1,6-trioxo-7-(2-pyridylmethylene)-2H,7aH-azetidino[2,1-b]1,3-thiazine-4-carboxylic acid (Biorg. Med. Chem. Lett. 2000, 853 and Biorg. Med. Chem. Lett. 2000, 847) and 3-(acetyloxymethyl)-7-[[tert-butyl]oxycarbonyl]methylene]-1,1,6-

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trioxo-2H,7aH-azetidino[2,1-b]1,3-thiazine-4-carboxylic acid (J. Med. Chem. 1995, 38, 1022).

b7) Carbapenem Derivatives of the Above General Formula IX:

Here, preferred combinations for Rk and Rl are: Rk=hydrogen and Rl=sulfamoyl or 3-carboxyphenyl (or vice versa); and Rk=Rl=methyl.

Preferred examples these compounds are imipenem, meropenem, ertapenem and doripenem.

b8) Boronate Derivatives of the Above General Formula X.

b9) Boronate Derivatives of the Above General Formula XI:

Examples of these compounds are 3-[(4-phenylsulfonyl-2-thienylsulfonyl)amino]phenyl]boronic acid (Chem. Biol. 2001, 8, 593) and [3-[[4-[(4-carboxyphenylsulfonyl)amino]phenylsulfonyl]amino]phenyl]boronic acid (Chem. Biol. 2001, 8, 594).

b10) Phosphonate Derivatives of the Above General Formula XII

Preferred examples of these are {[[(4-nitrophenoxy)(hydroxyphosphoryl)]methyl}[benzylsulfonyl]amine, {[[(4-nitrophenoxy)(hydroxyphosphoryl)]methyl}(phenylsulfonyl)amine and {[[(4-nitrophenoxy)(hydroxyphosphoryl)]methyl}(2-thienylsulfonyl)amine, (benzo[b]thiophen-2-ylsulfonyl){[(4-nitrophenoxy)(hydroxyphosphoryl)]methyl}amine and 2-[(4-nitrophenoxy)(hydroxyphosphoryl)]-1-(phenylsulfonyl)hydrazine (all described in US-A-2004/082546 and US-A-2004/029836); and

b11) Diazabicyclooctane Derivatives of the Above General Formula XIII:

A preferred example of these is (1R,2S,5R) 2-(aminocarbonyl)-7-oxo-1,6-diazabicyclo[3.2.1]octane-6-sulfonic acid (as described in WO-A-2002/01219, WO-A-2002/010172, FR-A-2835186 and FR-A-2848210).

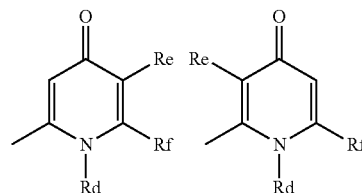
The pharmaceutical compositions of the present invention may comprise, besides the compound of formula I, two or more compounds selected from the above formulae II to XIII, being different from each other. Pharmaceutical compositions comprising triple combinations of a compound of formula I and two different compounds selected from the groups b1) to b11) are a preferred embodiment of the invention.

Preferred combinations for the pharmaceutical compositions of the invention are compounds of formula I, where R1 is SO₃H or OSO₃H or OCRaRa'COOH, wherein Ra and Ra' are as defined for formula I;

R2, R3 and R5 are as defined for formula I;

R4 is C(Rx)(Ry)Z, with either:

Rx=Ry=H and Z is a group of one of the formulae



wherein Rd, Re and Rf are as defined for formula I; or

Rx=Ry=methyl and Z=COOH;

or R4 is 4-hydroxypyridino[3,2-b]pyridin-3-yl)methoxy; and R6 is 2-amino-1,3-thiazol-4-yl, 2-amino-5-chloro-1,3-thiazol-4-yl, 5-amino-1,2,4-thiadiazol-3-yl, 5-aminopyrazol-3-yl or 4-aminopyrimidin-2-yl;

with either one of

(1S,5R)-2-[2-(3-carbamoylpyridyl)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, (1S,5R)-2-[2-(4-

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aminopyridyl)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, (1S,5R)-2-[2-(3-carbamoyl-6-methylpyridyl)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, (1S,5R)-2-[2-(5-methyl(1,3,4-thiadiazol-2-ylthio))acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, sodium salt, (1S,5R)-2-[2-(1-methyl(1,2,3,4-tetraazol-5-ylthio))acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, sodium salt, (1S,5R)-2-[N-[4-({[2-(2-aminoethoxy)ethyl]amino}carbonylamino)phenyl]carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid, (1S,5R)-2-[N-(4-{{[2-(2-aminoethyl)amino]carbonylamino}phenyl}carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid or (2S,3S,5R)-3-((1Z)-2-cyanovinyl)-3-methyl-4,4,7-trioxo-4-thia-1-azabicyclo[3.2.0]heptane-2-carboxylic acid.

More preferred are composition wherein the above double combinations are further combined with one of sulbactam, clavulanic acid or a customary pharmaceutically acceptable salt thereof, i.e. a clavulanate. These compositions thus comprise triple combinations.

Particularly preferred are compositions with the following double and triple combinations (the numbers are as in the foregoing tables 1a, 1b, 2a and 2b):

Aztreonam	Compound 102	Sulbactam
Aztreonam	Compound 102	Clavulanate
Compound 1	Compound 501	
Compound 1	Compound 102	Sulbactam
Compound 1	Compound 102	Clavulanate
Compound 1	Compound 103	Clavulanate
Compound 1	Compound 111	Clavulanate
Compound 1	Compound 202	Clavulanate
Compound 1	Compound 206	Clavulanate
Compound 1	Compound 323	Sulbactam
Compound 1	Compound 323	Clavulanate
Compound 1	Compound 324	Clavulanate
Compound 12	Compound 102	Sulbactam
Compound 12	Compound 102	Clavulanate
Compound 12	Compound 103	Sulbactam
Compound 12	Compound 323	Sulbactam
Compound 12	Compound 324	Sulbactam
Compound 21	Compound 102	Sulbactam
Compound 21	Compound 102	Clavulanate
Compound 21	Compound 323	Sulbactam
Compound 21	Compound 323	Clavulanate
Compound 21	Compound 324	Sulbactam
Compound 22	Compound 102	Clavulanate
Compound 22	Compound 324	Clavulanate
Compound 22	Compound 324	Sulbactam
Compound 26	Compound 102	Clavulanate
Compound 26	Compound 102	Sulbactam
Compound 26	Compound 324	Clavulanate
Compound 26	Compound 324	Sulbactam
Compound 29	Compound 102	Sulbactam
Compound 29	Compound 102	Clavulanate
Compound 29	Compound 323	Sulbactam
Compound 29	Compound 324	Sulbactam

Pharmaceutical compositions with triple combinations of compound no. 1 according to formula I with any one of the compounds of formula II and with clavulanate form a particularly preferred embodiment of the invention, such as do kits-of-parts (articles) with this combination.

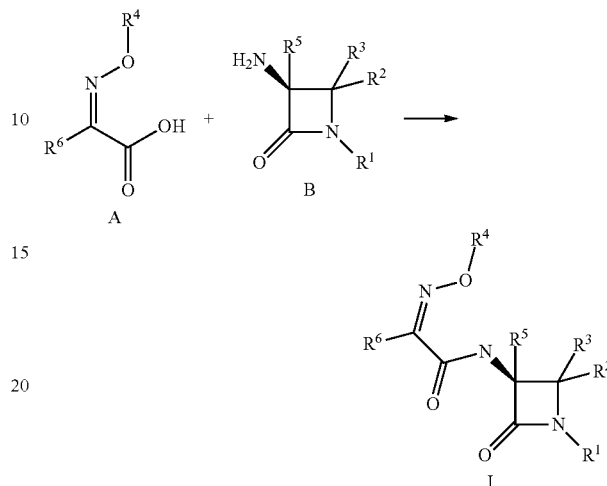
The compounds of formula I are compounds known from the above cited literature references, or can be made in an analogous manner, or can be made as described in the following. If in the following schemes a number is assigned to an intermediate then this intermediate is per se presumed novel and may form part of the invention.

The compounds of formula I can generally be prepared by reacting aryl or heteroaryl carboxylic acids of general formula A with 3-amino-azetidin-2-one compounds of general

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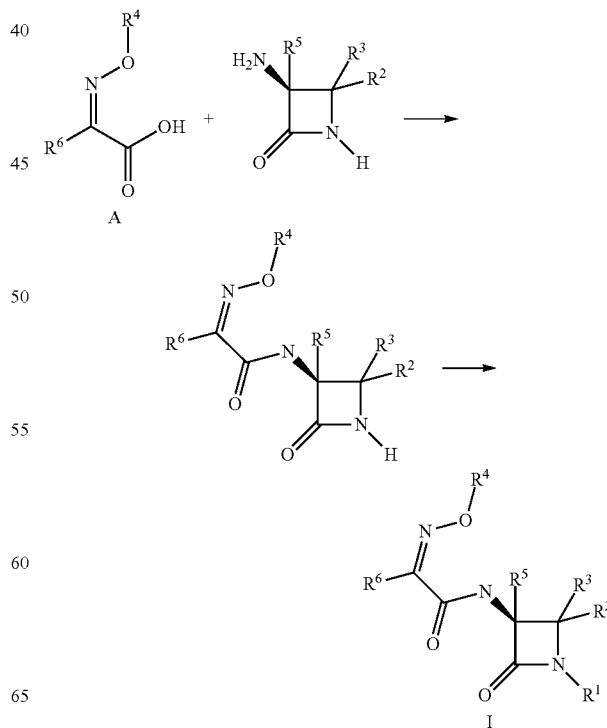
formula B (scheme 1). In this scheme, R4 may also have the meaning of a protecting group, which may then be removed, in order to subsequently connect the actually desired R4.

Scheme 1



The coupling reaction of compounds of general formula A with compounds of general formula B can be performed with the corresponding acyl chlorides of the aryl or heteroaryl carboxylic acids of general formula A or with the carboxylic acids A themselves and DCC (Chem. Pharm. Bull 1983, 2200) or with an activated ester of the aryl or heteroaryl carboxylic acids of general formula A, such as the N-hydroxysuccinimidyl ester (see Org. Process Res. & Dev. 2002, 863) or the benzothiazolyl thioester (see J. Antibiotics 2000, 1071). Alternatively, compounds of formula I can also be prepared as outlined in scheme (with R¹=SO₃H see also J. Antibiotics 1985, 346; J. Org. Chem. 1981, 1557).

Scheme 2

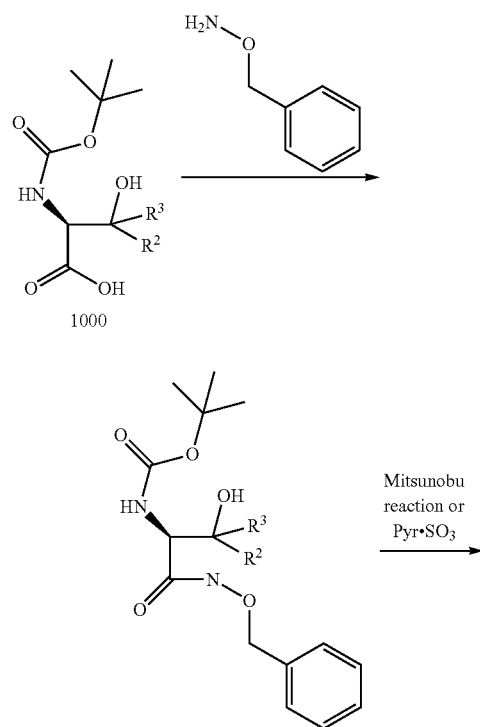


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A) Preparation of Compounds B

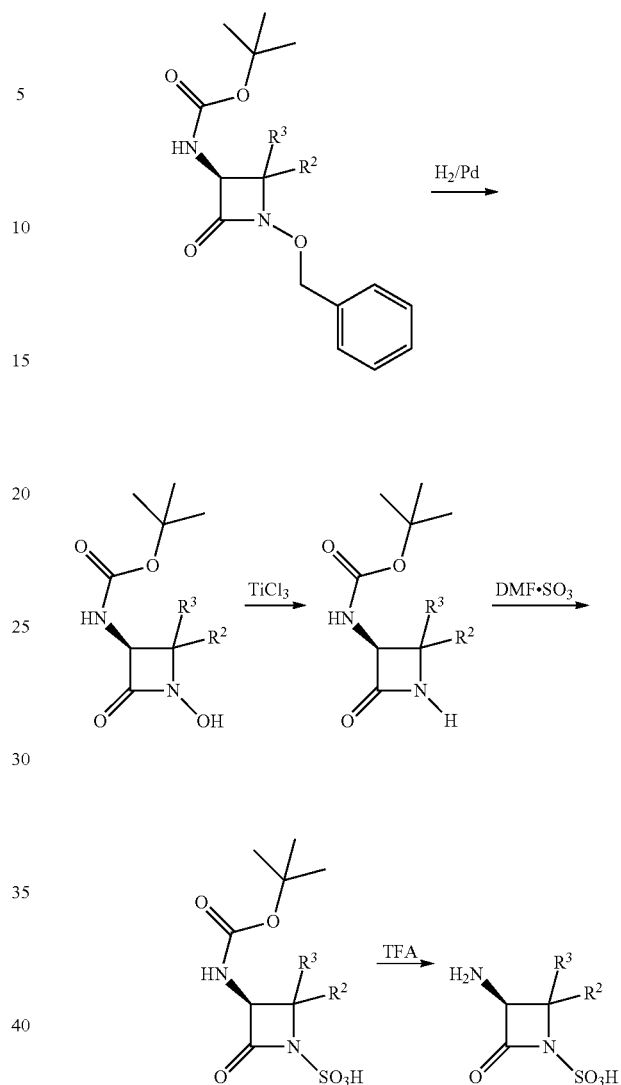
The preparation of compounds of general formula B can be carried out in different ways according to the substituents present in 1, 3 and 4-positions (Scheme 3, 4, 5). (J. Org. Chem. 1980, 410; J. Org. Chem. 1985, 3462; J. Antibiotics 1985, 346; J. Antibiotics 1985, 813; J. Antibiotics 1986, 76; Tetrahedron Lett. 1986, 2789, J. Med. Chem. 1985, 1447; Chem. Pharm. Bull. 1984, 2646, J. Am. Chem. Soc. 1990, 760).

Scheme 3

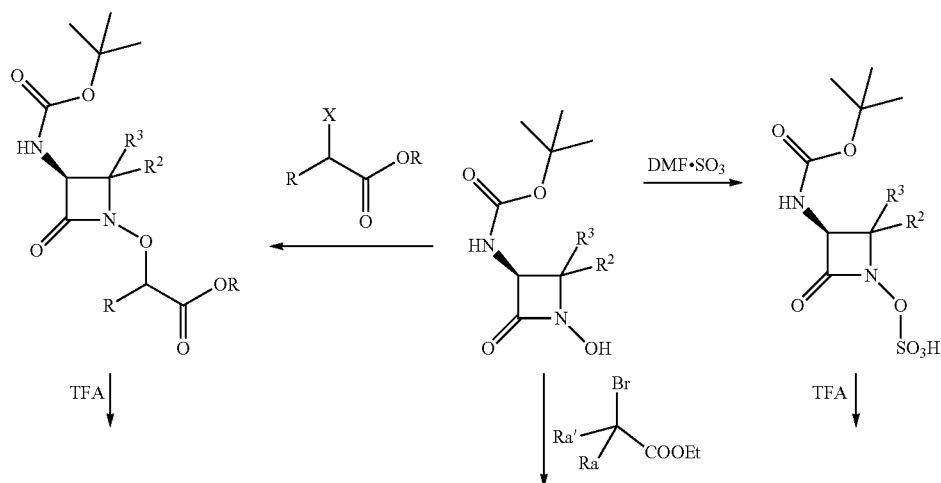


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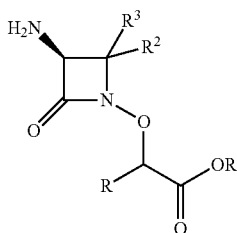
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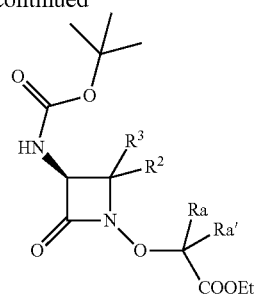
Scheme 4



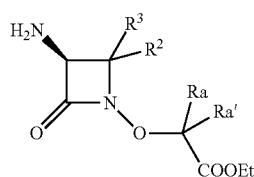
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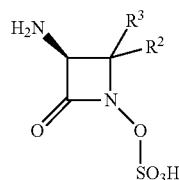
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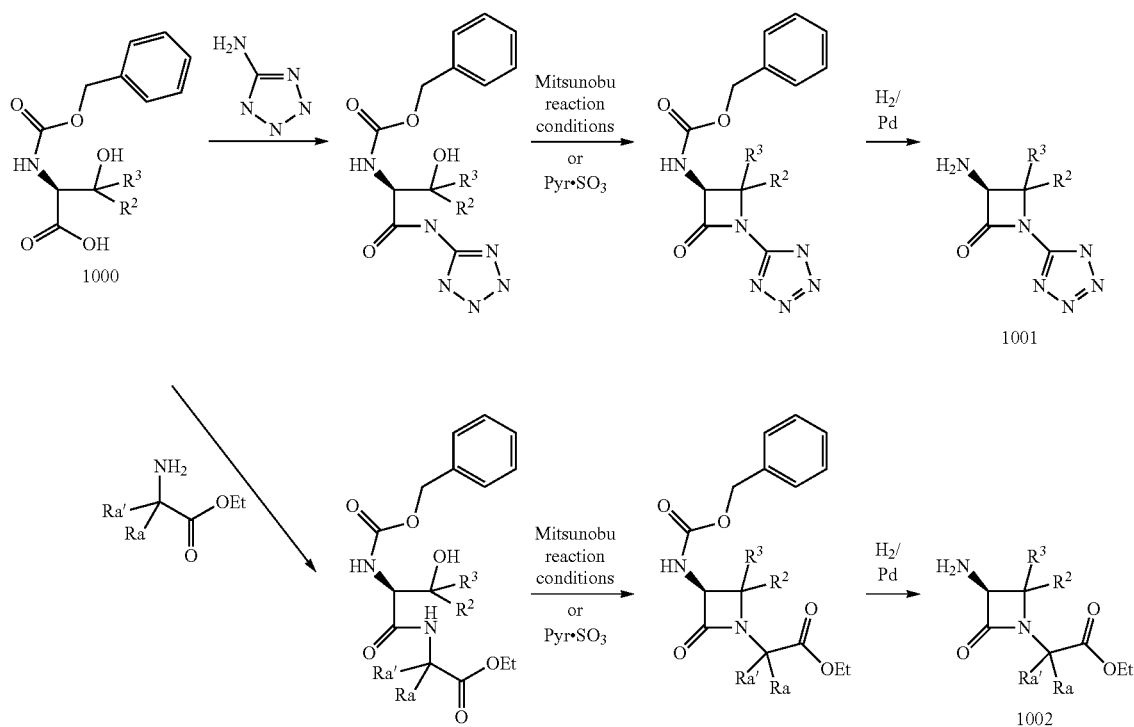
TFA



48



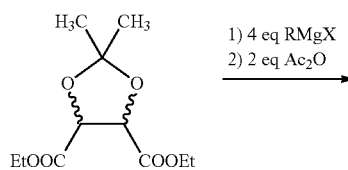
Scheme 5

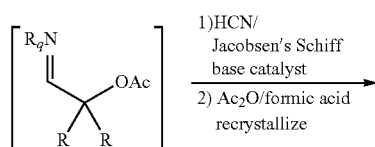
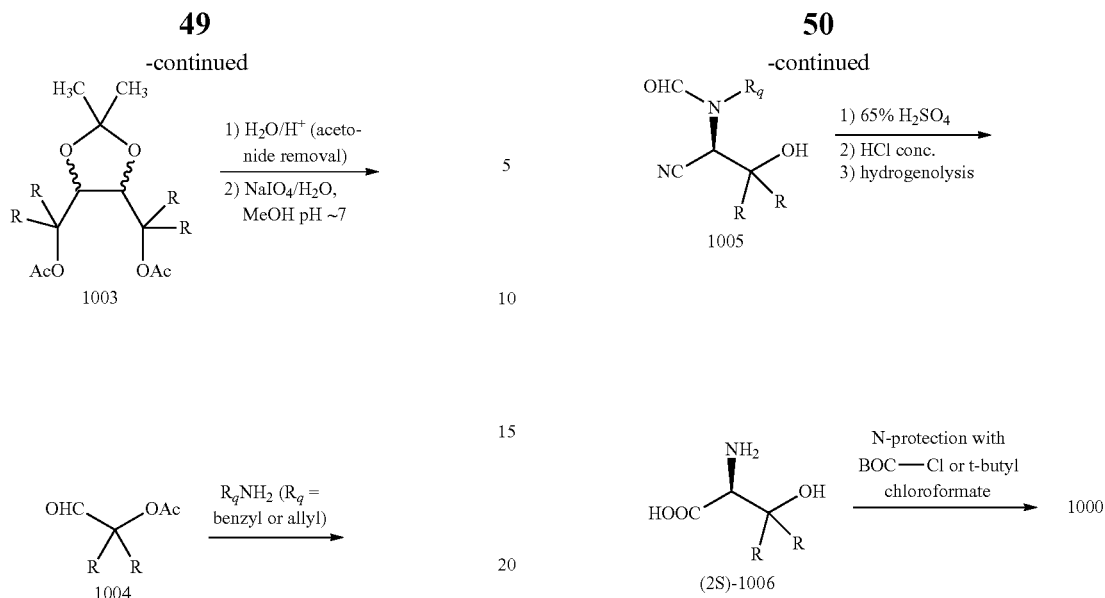


A-1) In the above schemes 3 and 5, generally an enantiomerically pure N-protected β -hydroxy amino acid 1000 is required as the starting material. This precursor 1000 can be prepared in different ways, as outlined in the following sections A-1-I) to A-1-IV):

A-1-I) Where $R^2=R^3=R$ and preferably is alkyl, alkenyl, alkynyl, optionally substituted benzyl or optionally substituted phenyl, a synthesis according to the following scheme A-1-I or scheme A-1-Ia may be used:

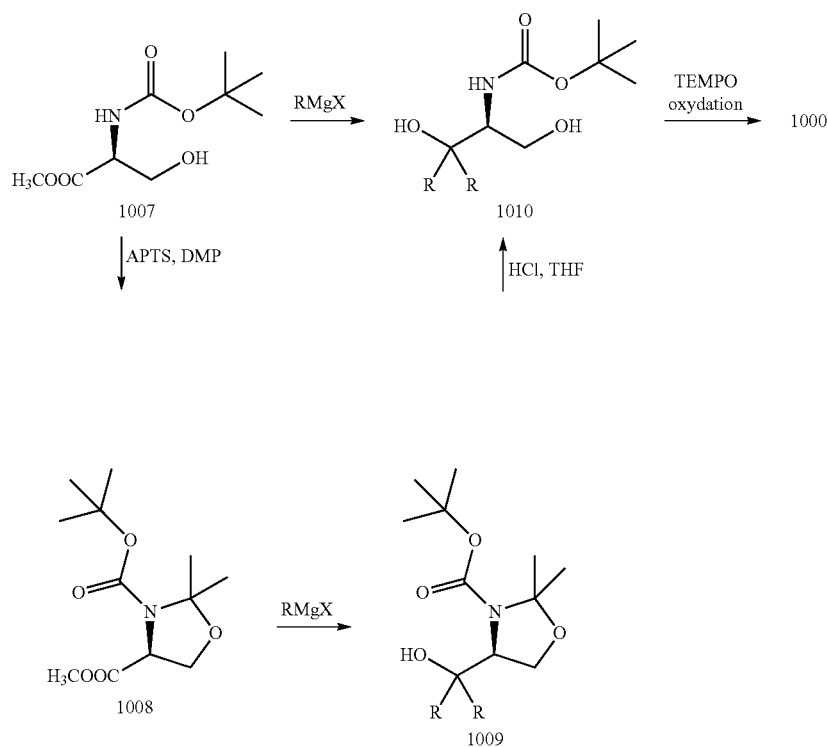
Scheme A-1-I





In the above scheme A-1-I, the configuration of the α -carbon atoms in the starting diethyl tartrate needs not be defined. The indicated chiral Jacobsen's catalyst for the asymmetric Strecker synthesis of the amino acid (2S)-1006 has been known from Angew. Chem. Int. Ed. 2000 1279. As the catalyst is available in both enantiomers, the corresponding β -hydroxy amino acid with the D-configuration at the α carbon atom ((2R)-1006) may also be produced. The latter is suited for producing N-protected β hydroxy amino acids 1000a where the hydrogen atom at the α carbon atom is replaced by R5 (see section A-1-IV below).

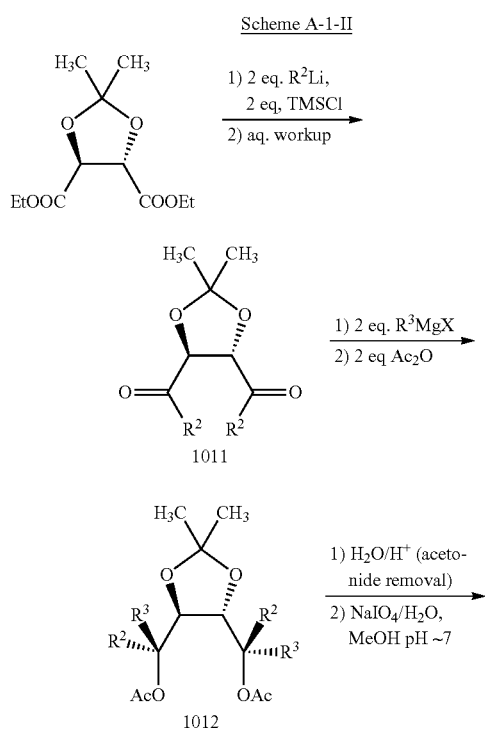
Scheme A-1-Ia



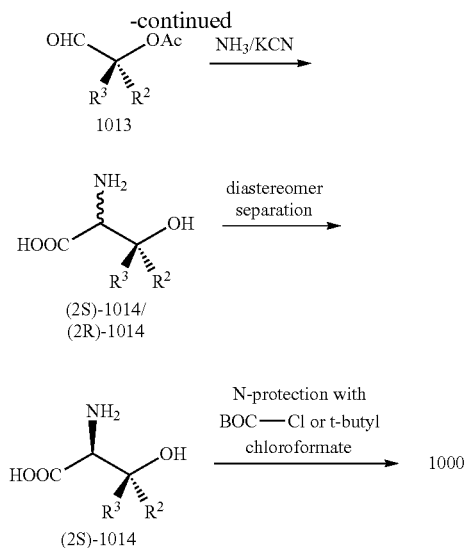
51

In the above scheme the preparation of the β,β -dialkylsubstituted β -hydroxy α -amino acid 1000 starts from the commercially available enantiomerically pure N-BOC serine methyl ester 1007 (J. Org. Chem. 2003, 177, Tetrahedron, 1996, 11673). The synthesis follows the chemistry based on Rapoport's methodology, which is known to keep the conformational integrity of the starting amino acid (J. Org. Chem. 1989, 1866, J. Org. Chem. 1990, 3511).

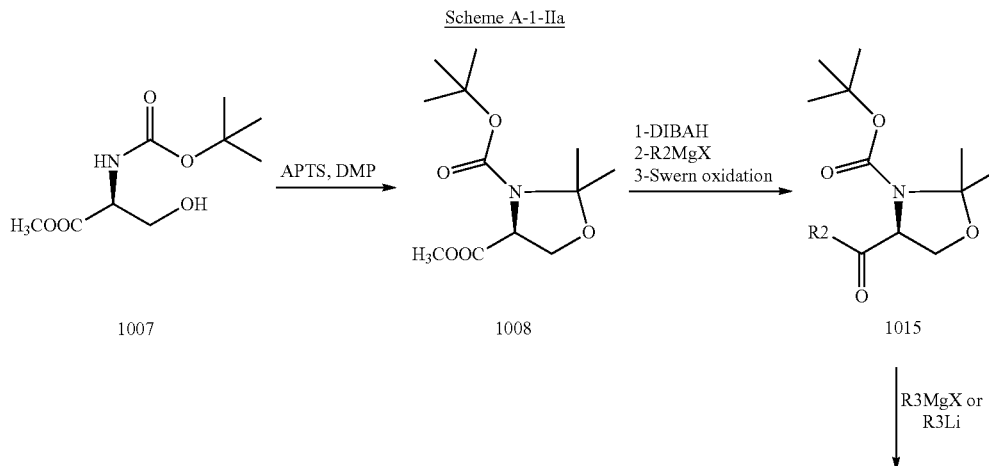
A-1-II) Similarly, where R^2 and R^3 are different from each other and are preferably independently selected from alkyl, alkenyl, alkynyl, optionally substituted benzyl and optionally substituted phenyl, the following scheme A-1-II, modified from scheme A-1-I, may be used to produce 1000:



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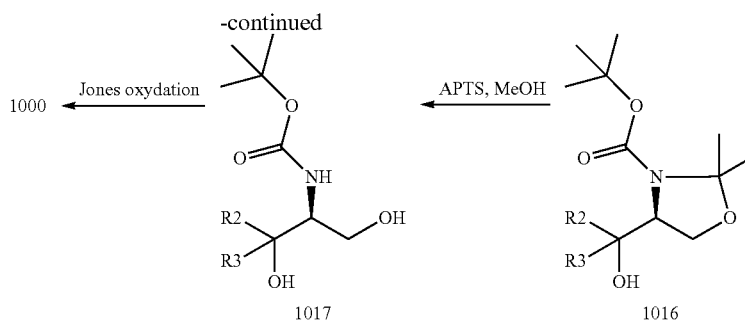


In this scheme R^2 is particularly preferably methyl. The introduction of the second residue R^3 into 1011 by Grignard reaction gives, according to Cram, predominantly the shown diastereomer of 1012, due to the chelating effect of the oxygen atom of the α -acetonide substituent. In the Strecker synthesis step without chiral auxiliary, both diastereomeric amino acids (2S)-1014 and (2R)-1014 may form as a mixture due to the newly formed chiral α -carbon atoms. The diastereomers of the so produced β -hydroxy amino acids 1014 may be separated using ion exchange chromatography with aqueous buffers as the mobile phase, as is customary in the art. The correct diastereomer (2S)-1014 may be identified as the one that produces faster a dipeptide, when each of the diastereomers (2S)-1014 and (2R)-1014 is reacted under otherwise identical conditions with N-benzoyl L-alanine methyl ester and carboxypeptidase Y as the dipeptide-forming enzyme (for an appropriate experimental procedure see example 1 of EP-A-0 017 485). The other, undesired diastereomer (2R)-1014 may be used for the production of N-protected β -hydroxy amino acids 1000a where the hydrogen atom at the α carbon atom is replaced by R^5 other than hydrogen (see section A-1-IV below).



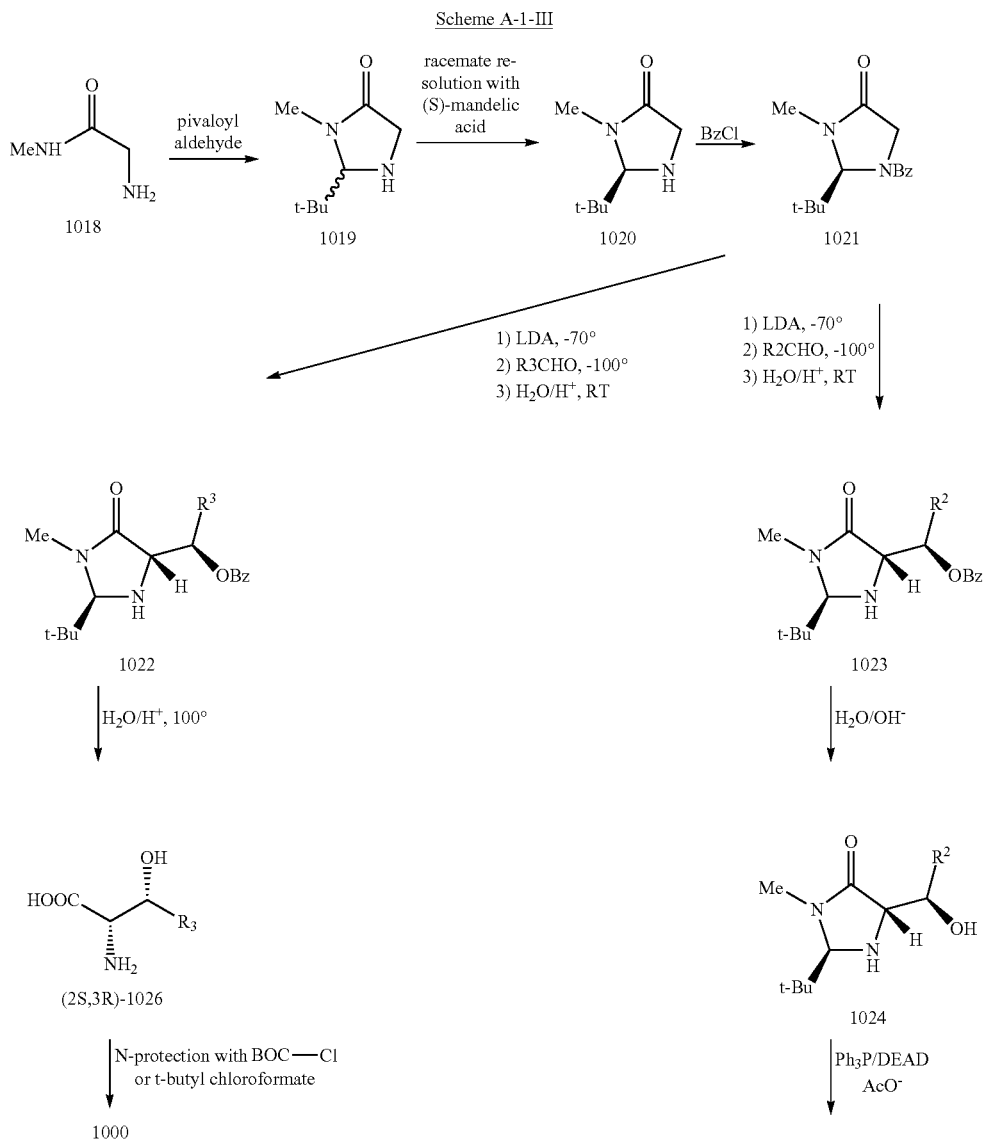
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Scheme A-1-IIa outlines an alternative procedure starting from protected serine 1007 and leading to disubstituted hydroxyl derivatives 1016 by controlling the stereochemistry of the addition of the second substituent; either with R^3Li , giving the Felkin adduct as major product, or with R^3MgX , to obtain the anti-Felkin adduct as major product (Tetrahedron 1995, 8121).

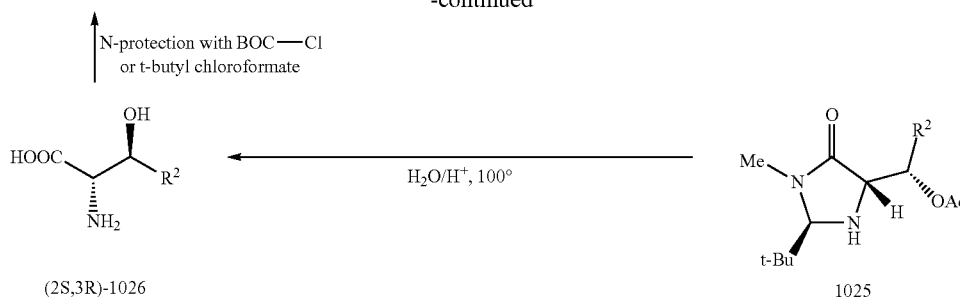
A-1-III) where R^2 is preferably alkyl, alkenyl, alkynyl, optionally substituted benzyl or optionally substituted phenyl, and R^3 is H, or R^2 is H and R^3 is preferably alkyl, alkenyl, alkynyl, optionally substituted benzyl or optionally substituted phenyl, a synthesis according to following scheme A-1-III) may be adopted to form 1000:



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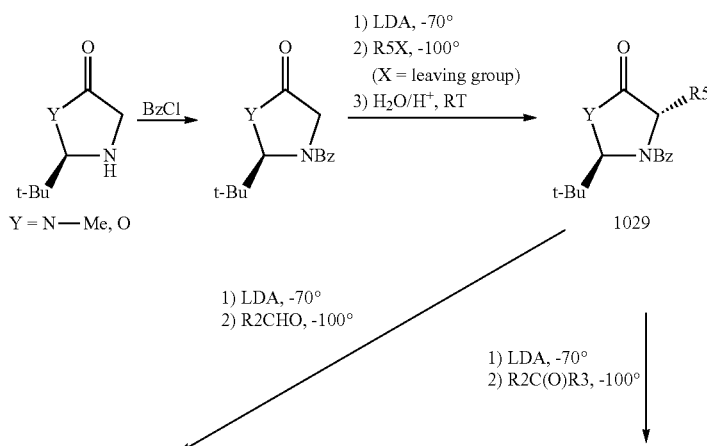
This methodology was developed by Seebach (Helv. Chim. Acta 1987. 237). In scheme A-1-III, the conversions on the left pathway yield the N-protected β -hydroxy amino acid 1000 where R^2 =hydrogen, in defined configuration. The conversions on the right pathway yield the N-protected β -hydroxy amino acid 1000 where R^3 =hydrogen, also in defined configuration.

A-1-IV) When the azetidinone B contains R_5 other than hydrogen, then an N-protected α - R^4 -substituted β -hydroxy amino acid 1000a is required, analogous to a corresponding above amino acid 1000, except for the additional R_5 substituent. This amino acid 1000a may be used in above schemes 3 and 5 instead of 1000.

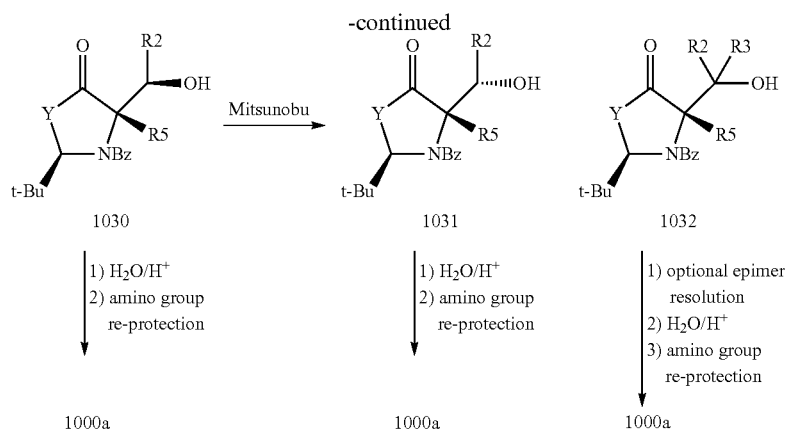
A-1-IVa) When R_5 is preferably alkyl, and one of R_2 and R_3 is preferably hydrogen, and the other one preferably is alkyl, alkenyl, alkynyl, optionally substituted phenyl or optionally substituted benzyl (or both of R_2 and R_3 are preferably independently selected from alkyl, alkenyl, alkynyl,

optionally substituted phenyl and optionally substituted benzyl) the technology described in part A-1-III (Helv. Chim. Acta 1987. 237) can be employed, but using 2-tert-butyl-N-benzoyl-1,3-oxazolidinone as chiral inductor (scheme A-1-IVa below). The introduction of the electrophile $\text{R}_5\text{-X}$ leads to the compound 1029. A condensation with aldehyde R_2CHO as second electrophile gives the R_2, R_5 -disubstituted oxazolidinone 1030 with control of the stereochemistry. If desired, the configuration at the newly formed secondary alcohol in 1030 may be inverted, such as under Mitsunobu conditions, to form the epimeric R_2, R_5 -disubstituted oxazolidinone 1031. If onto 1029a condensation with ketone $\text{R}_2\text{C}(\text{O})\text{R}_3$ as the second electrophile is carried out, then $\text{R}_2, \text{R}_3, \text{R}_5$ -trisubstituted oxazolidinones 1032 may be produced. If necessary the formed epimers of 1032 may be separated, such as by chromatography. All three compounds 1030, 1031 and 1032 may subsequently be converted by hydrolysis of the benzoyl group and the oxazolidinone ring and re-protection of the amino group desired compounds 1000a.

Scheme A-1-IVa



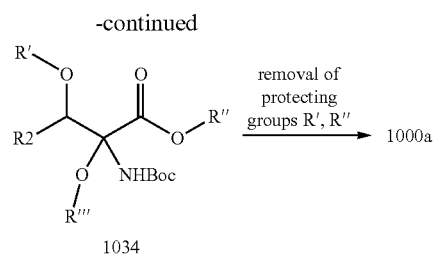
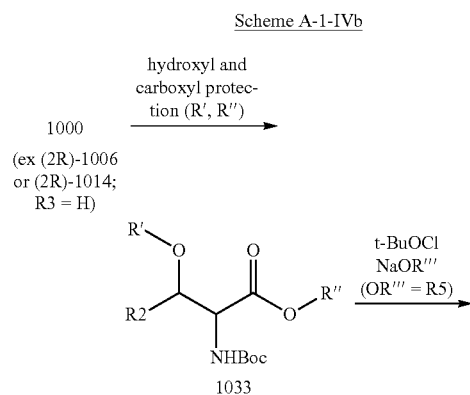
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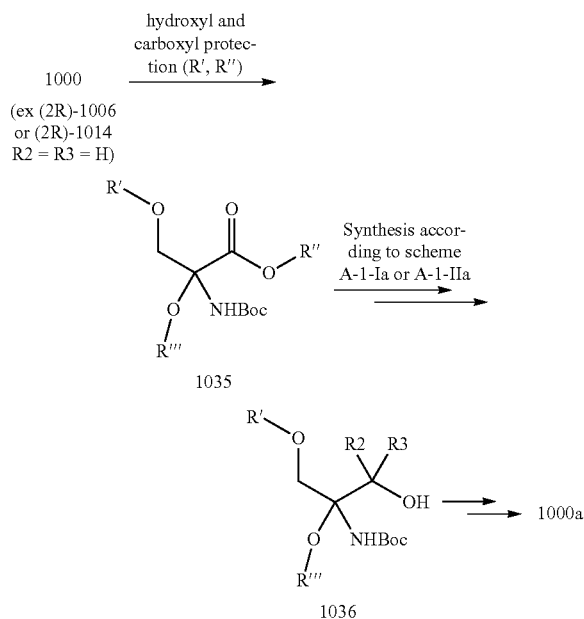
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A-1-IVb) When the azetidinone B contains alkoxy as R5 and one of R3 or R2 preferably as hydrogen, and the other, one preferably as alkyl, alkenyl, alkynyl, optionally substituted phenyl or optionally substituted benzyl, then another example of preparation of amino acids 1000a is outlined in following scheme A-1-IVb (in the scheme R3 is assumed as hydrogen). It is based on the chemical description written on Biochemistry 2004, 3385 and Fortsch. Chem. Org. Naturst., 1979, 327. The approach relies on the established oxidation of an N-acyl α -amino ester to a highly reactive intermediate N-acyl α -imino ester, which then adds R5-containing nucleophiles (alcohols R'''OH wherein R'''O=R5). The various protecting groups R' and R'' would be removed subsequently (not shown in the scheme).

A-1-IVc) Similarly, for the case where azetidinone B contains alkoxy as R5 and both R3 and R2 are preferably independently selected from alkyl, alkenyl, alkynyl, optionally substituted phenyl and optionally substituted benzyl, an approach according to following scheme A-1-IVc may be used. By this scheme it is possible to prepare β -disubstituted N-protected amino acids 1000a following the synthetic pathway described in scheme A-1-Ia and scheme A-1-IIa.



Scheme A-1-IVc

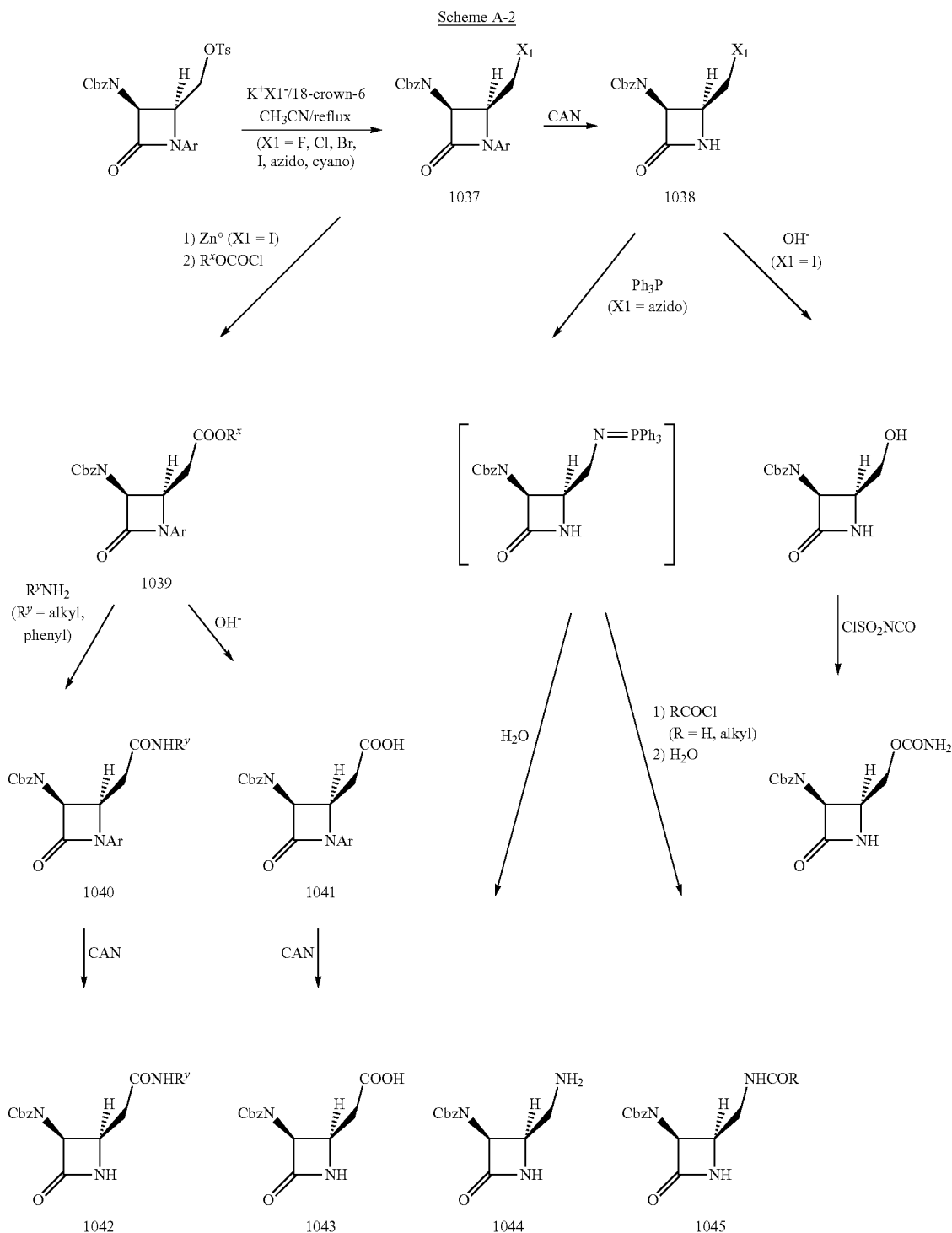


In above scheme A-1-IVc, the conversions 1000 to 1035 are analogous to the conversions of 1000 to 1034 in above scheme A-1-IVb.

A-2) A range of compounds B, where R2=CH₂X1 and R3=H, can be directly made in analogy to the procedure described in J. Antibiotics, 1983 1201 followed by standard functional group conversions (scheme A-2):

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This scheme shows how to form compounds B where X1 is preferably selected from halogen, azido, amino, hydroxyl, cyano, carboxyl, alkoxycarbonyl, alkanoylamino, phenylaminocarbonyl, alkylaminocarbonyl, aminocarbonyl, carbamoyloxy, alkylaminosulfonyl and optionally substituted phenylaminosulfonyl. The common starting material shown in the upper left of scheme A-2 is known from J. Org. Chem. 1982, 2765. In this scheme, in all the obtained compounds B

the final removal of the Cbz protecting group by hydro-

60 genolysis is not shown.

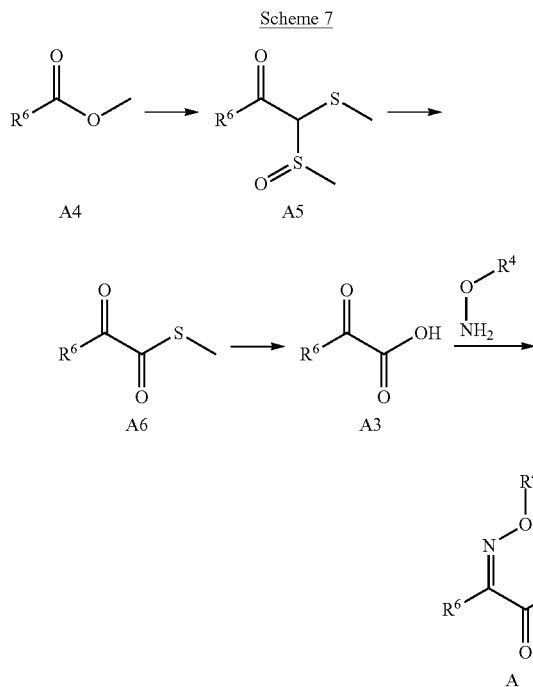
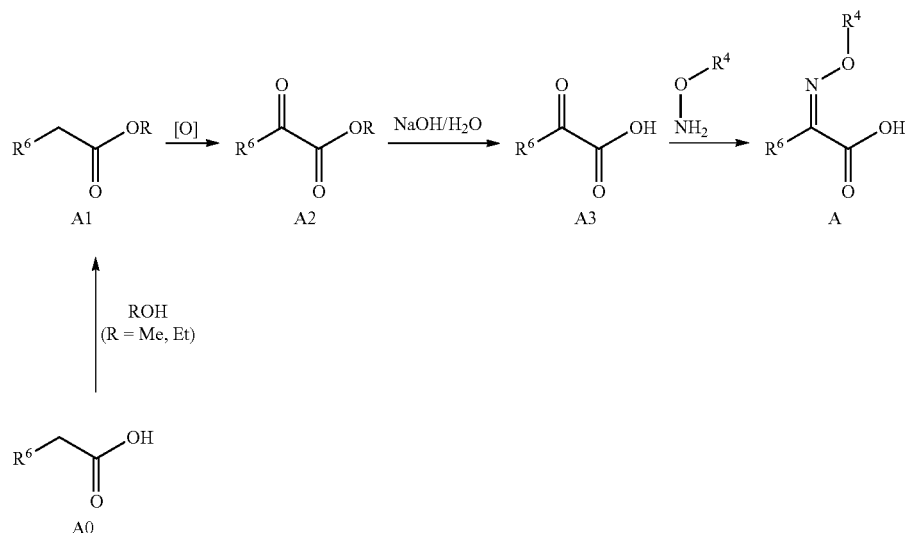
B) Preparation of Compounds A

The preparation of compounds of general formula A required in schemes 1 and 2 can be carried out in a customary way by reacting an appropriately R6-substituted keto acid A3 with an appropriately R4-etherified hydroxylamine (schemes 6 and 7 below)

61

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Scheme 6



The R⁶-substituted keto acid A₃ required in both schemes 6 and 7 can be prepared via 2 different synthetic pathways, as described in the following.

A₃ may firstly be obtained by oxidation of an ester A₁ leading to the glyoxalate derivative A₂, followed by hydrolysis of the ester group (scheme 6). The oxidising agent used in the conversion from A₁ to A₂ is not critical. Examples of suited oxidising agents are SeO₂ (J. Antibiotics 1983, p. 1020ff.), DMSO, X₂ (Bull. Chem. Soc. Jpn. 1994, 1701), X₂ and pyridine-N-oxide (Bioorg. Med. Chem. 2003, 591) where X₂ is a halogen; with SeO₂ being preferred.

The R⁶-substituted keto acid A₃ can also be prepared via the condensation of methyl methylthiomethyl sulfoxide

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according to above scheme 7 (J. Antibiotics, 1984, 546, J. Antibiotics, 1984, 557) in a 4 step synthesis from the esters A₄ R⁶COOCH₃ or R⁶COOEt. The methyl methylthiomethylsulfoxide is first condensed with A₄ derivatives to give the methyl thioglyoxylate compound A₆ after acidic treatment.

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Examples of preparation of intermediates A₀, A₁ or A₄ (scheme 6 and 7) are as outlined in the following subsections B-1-Ia) to B-2-Id). Some of these intermediates are also commercially available.

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B-1-I) R₆ can be a 5 membered heteroaromatic ring containing 1 to 4 heteroatoms such as N, O, S and which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, carbonylamino and halogen such as F, Cl, Br or I, preferably Cl.

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B-1-Ia) Thiadiazole Derivatives

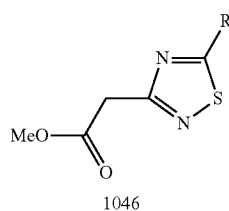
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The preparation of derivatives A₀, A₁ or A₄, where R₆ is a thiadiazole, especially 1,2,4-thiadiazole, may be performed as described in Biorg. Med. Chem. 2006, 1644, included herein by reference. Examples of compounds A₁, may be prepared starting from 3-amino-5-methoxyisoxazole or methyl amidine in presence of thioesters, potassium thiocyanate or isothiocyanate as outlined in scheme B-1-Ia (Bull Chem Soc Jpn. 1994, 1701, J. Antibiotics 1983, 1020). In this scheme, R is preferably selected from hydrogen, alkyl, alkoxy, amino, alkylamino and carbonylamino.

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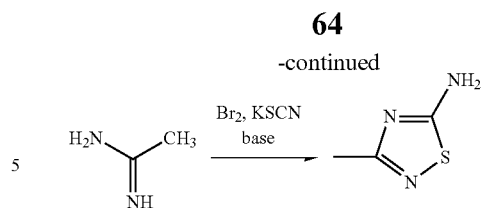
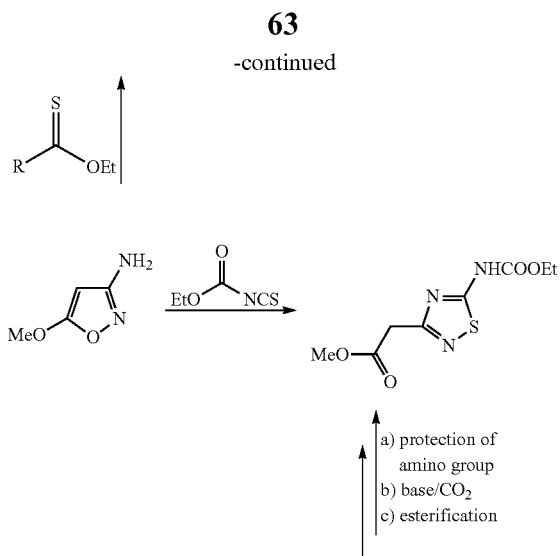
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Scheme B-1-Ia



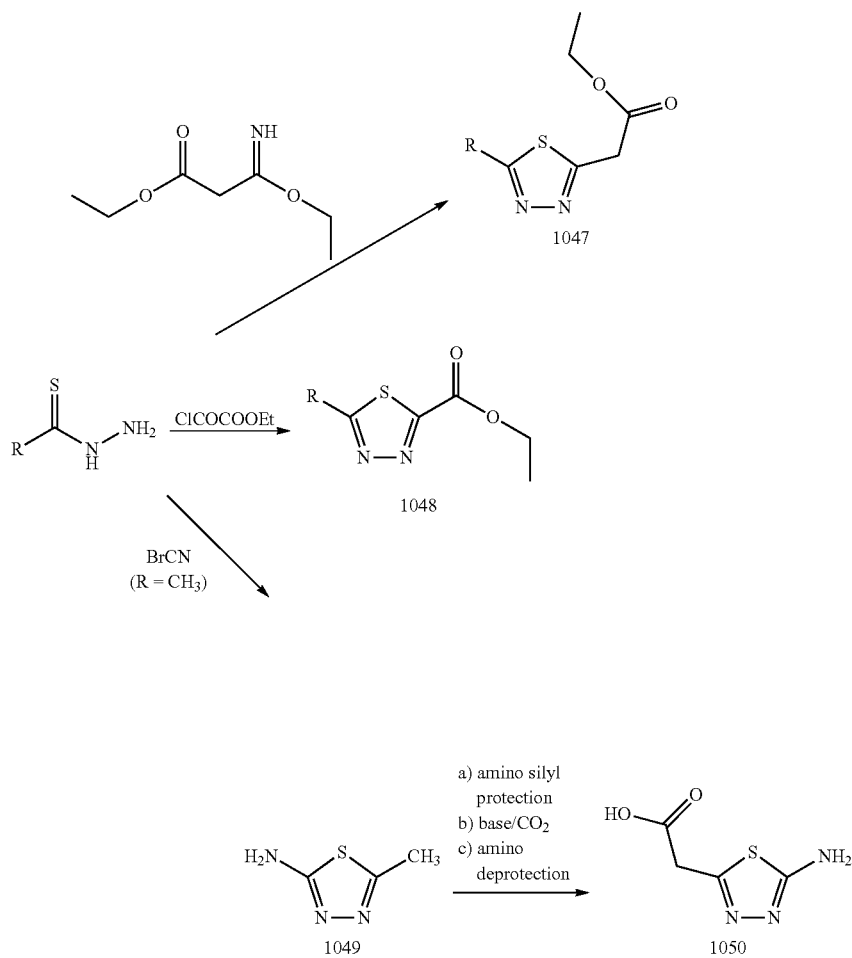
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B-1-Ib) Scheme B-1-Ib outlines examples of starting materials. A0, A1 and A4 (Helv. Chim. Acta 1982, 2606; Russ. J. Org. Chem. 2003; 1133, Tetrahedron Let. 1979, 2827) obtained from substituted thiohydrazines. Using BrCN and these as starting material, 1049 may be prepared and then can lead to 2-amino(1,3,4-thiadiazol-5-yl)acetic acid 1050. In scheme B-1-Ib, R is preferably selected from hydrogen, alkyl, alkoxy, amino, alkylamino and carbonylamino.

Scheme B-1-Ib

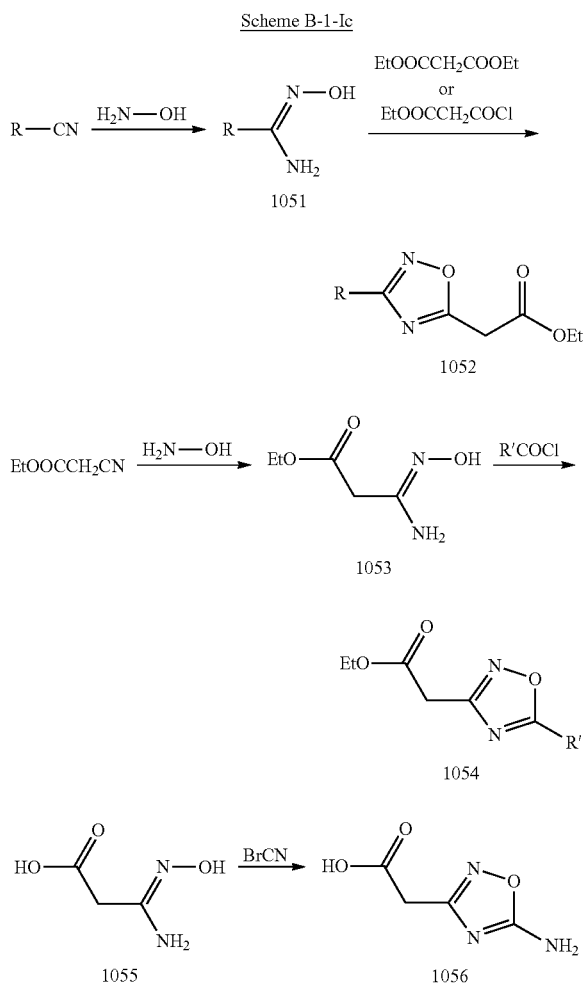


In the above schemes B-1-Ia and B-1-Ib, the reaction with base/ CO_2 , optionally by silyl protection of the amino group, may be carried out according to J. Antibiotics, 1984, 532 already cited above.

65

B-1-Ic) 1,2,4-oxadiazole Derivatives

As outlined in the upper part of scheme B-1-Ic indicated below, amide oxime derivatives 1051, wherein R is preferably selected from hydrogen, alkyl and alkoxy may be reacted with compounds such as carboxylic acid, acyl chloride or cyano derivatives to form 1,2,4-oxadiazole rings 1052, which are new examples of heterocycles derivatives A1. The lower part of scheme B-1-Ic, wherein R' is preferably selected from hydrogen, alkyl, alkoxy, carbonylamino and halogen (in particular Cl), shows the synthesis of carboxylates 1054, 1056 of type A1 which are isomeric to 1052 (Tetrahedron Lett. 1998, 3931, J. Org. Chem 1995, 3112, J. Med. Chem. 1990, 1128, J. Pep. Research 2003, 233, Z. Chem. 1975, 57). For instance, compound 1056 may be obtained from the condensation of 1055 with BrCN.

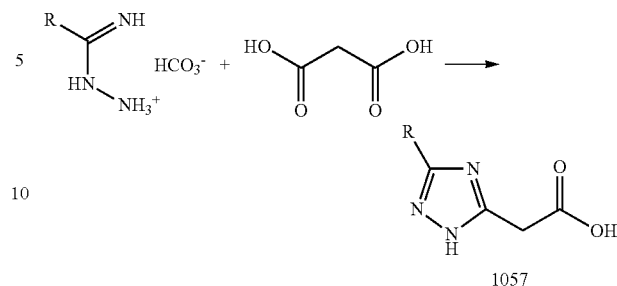


B-1-Id) Triazole Derivatives

Other examples of acids A0 outlined in scheme B-1-Id shown below are triazolyl acetic acid derivatives, such as 1,2,4-triazolyl acetic acids 1057, which may optionally be substituted by one substituent R preferably selected from amino, alkyl, alkoxy and carbonylamino. The exemplary synthesis of 2-(5-amino-1,2,4-triazol-3-yl)acetic acid (R=NH₂) by this scheme is known from Russ. J. Org. Chem. 1995, 240.

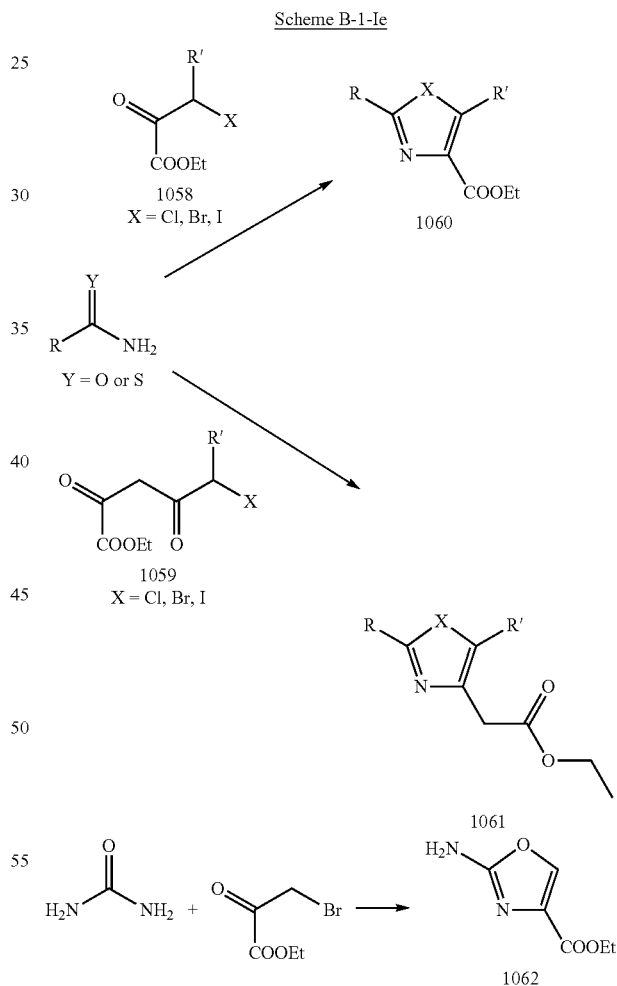
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Scheme B-1-Id



B-1-Ie) Thiazole and Oxazole Derivatives

A versatile way for forming intermediates A0, A1 or A4, wherein the residue R6 is an optionally substituted 1,3-thiazol-4-yl or 1,3-oxazol-4-yl, is according to the known reactions of a thioamide or amide with an α -haloketone derivatives, as outlined in scheme B-1-Ie):



In this scheme, R may preferably be selected from hydrogen, alkyl, alkoxy, amino, and alkylamino; and R' is preferably selected from hydrogen and alkyl. From substituted thioamides, and substituted ethyl halogeno pyruvate 1058 or compounds 1059 substituted 1,3-thiazol-4-yls 1060, 1061 may be obtained according to examples described in the

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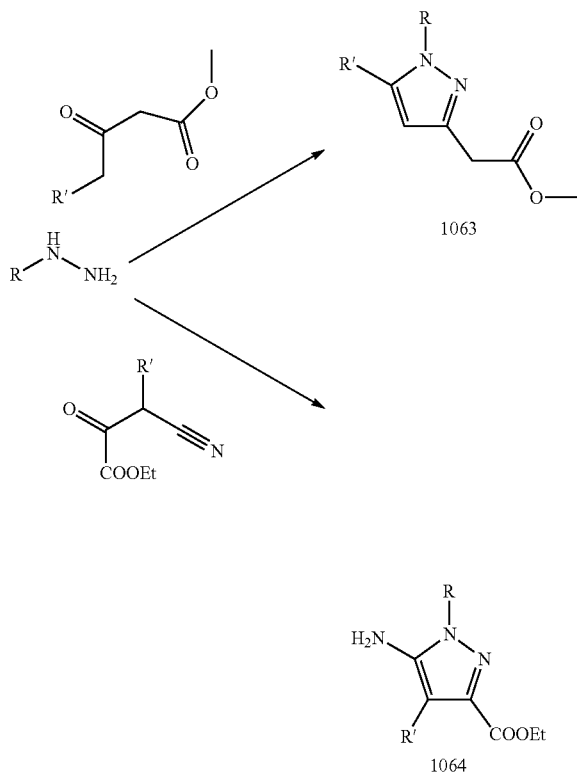
literature such as Tetrahedron Lett. 2005, 66; J. Chem. Soc. 1966, 1357; J. Chem. Soc. 1960, 925; J. Med. Chem. 1971, 1075; J. Het. Chem. 1980, 1255; J. Med. Pharm. Chem. 1959, 577.

From similar keto ester derivatives 1059 in the presence of amide or thioamide derivatives, substituted 1,3-oxazol-4-yl acetic acid esters may be prepared as reported for instance in Bioorg. Med. Chem, 2003, 4325; Heterocycles, 2001, 689; Chem. Pharm. Bull. 1986, 2840; Tetrahedron Lett. 1992, 1937. The preparation of compounds 1062 from urea with ethyl bromo pyruvate is an example of preparation of 1,3-oxazole derivatives. If R' is hydrogen and X is sulphur, then the thiazole moiety in 1060 or 1061 may subsequently be chlorinated using the procedure as described in "Preparation 1" of EP-A-0 055 465.

B-1-If) Pyrazole Derivatives.

Another example of heterocyclic intermediates A0, A1 or A4 may be synthesised from substituted ethyl pyruvates in presence of hydrazine or substituted hydrazines according to the scheme B-1-If (J. Chem. Soc. 1945, 114; Helv. Chim. Acta 1955, 670, J. Am. Chem. Soc. 1959, 2456). In this scheme, R is preferably selected from hydrogen, alkyl and carbonylamino, and R' is preferably selected from hydrogen, alkyl, alkoxy, carbonylamino, hydroxyl, amino, alkylamino and halogen (in particular Cl).

Scheme B-1-If



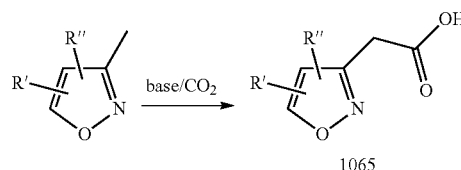
The reaction shown in the lower part of the scheme is analogous to a synthesis described in J. Org. Chem. 2004, 5168.

68

B-1-Ig) Isoxazole Derivatives

Many isoxazoles with a carboxyl substituent in the 3-position and with one or two substituents selected from amino, alkyl (in particular methyl and ethyl) and hydroxyl are commercially available. Similar commercially available isoxazoles with a methyl substituent in the 3-position and optionally one or two substituents R' and R'' preferably independently selected from alkyl, alkoxy and halogen, may again be converted to corresponding carboxylate-containing isoxazoles by converting that 3-methyl substituent to carboxylate using base and carbon dioxide (scheme B-1-Ig). For example, in U.S. Pat. No. 4,394,504 3-amino-5-isoxazolyl-2-acetic acid 1065 was prepared from 3-amino-5-methylisoxazole in this way.

Scheme B-1-Ig



B-2) R6 can also be a phenyl ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, carbonylamino and halogen or a 6-membered heteroaromatic ring containing 1 to 5 heteroatoms such as N and which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, carbonylamino and halogen (such as F, Cl, Br, I, preferably Cl).

B-2-Ia) A very general way of obtaining keto acids A3 for above schemes 6 and 7 is by direct Friedel-Crafts acylation of an appropriately substituted corresponding phenyl or heterocycle R6-H, using excess oxalyl chloride. This way is feasible for all keto acids A3 where R6 is phenyl or a heterocycle which may be substituted by one to five substituents selected from alkyl, alkoxy, dialkylamino and halogen (in particular chloro), with the proviso that the phenyl or heterocycle has at least one unsubstituted carbon atom (the carbon atom that will carry the glyoxaloyl substituent). The appropriately substituted corresponding phenyl or heterocycle R6-H (where H is bonded to the said unsubstituted carbon atom) is reacted with excess oxalyl chloride under Friedel-Crafts conditions, followed by removal of excess oxalyl chloride and hydrolysis of the remaining free acyl chloride group of the introduced glyoxaloyl moiety. Friedel-Crafts acylation is also feasible for such phenyls or heterocycles where some of the one to five substituents are hydroxyl, amino, alkylamino. These substituents, however, are preferably protected as the before the acylation step and the subsequently deprotected.

B-2-Ib) Examples of intermediates A4 are commercially available pyridinecarboxylic acids, such as picoline (2-pyridinecarboxylic acids), nicotinic acid (3-pyridinecarboxylic acids) or isonicotinic acid (3-pyridinecarboxylic acids), which may optionally be substituted at the pyridyl by a substituent selected from alkoxy, halogen (in particular chloro) and amino.

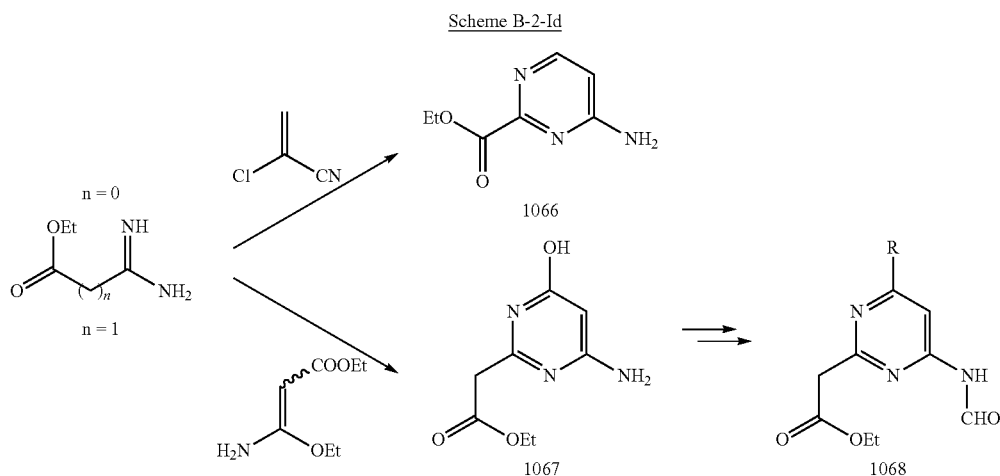
69

B-2-Ic) Further examples of acids A0 are 2-pyridyl acetic acids, such as 2-(pyridyl-2) acetic acids 2-(pyridyl-3) acetic acids or 2-(pyridyl-4) acetic acids which may optionally be substituted at the pyridyl by a substituent selected from alkoxy, halogen (in particular chloro) and amino.

These can be obtained by deprotonating an appropriately substituted methylpyridine with as strong base such as N-BuLi or LDA and reacting the anion with carbon dioxide. In this reaction, the methyl substituent of the methylpyridine is at the position where the acetic acid will be. The optional amino substituent at the pyridine may have been appropriately protected beforehand, such as with TMS-Cl. An exemplary reaction of this type can be found in DE-OS-2848912 and J. Antibiotics 1984, 532.

B-2-Id) R6 can also be a pyrimidine derivative.

Scheme B-2-Id (below) shows that amidine derivatives (J. Org. Chem. 1962, 3608) can lead to either 2-substituted pyrimidine A1 or A4 (DE-OS-2848912, J. Antibiotics 1984, 546). Compounds 1068 (R=Cl) can easily be obtained by reaction with phosphoryl chloride from 1067. The chlorine can then easily be substituted by nucleophiles such as ammonia, alkylamines or alcohols and lead to compounds 1068 with R=amino, alkylamino or alkoxy (J. Antibiotics 1984, 546).



Another group of commercially available or synthetically accessible esters A4 are pyrimidyl-4 (ethyl 5,6-diaminopyrimidine-4-carboxylate, ethyl 2-amino-5-chloropyrimidine-4-carboxylate), pyrimidyl-5 (ethyl 2,4-diaminopyrimidine-5-

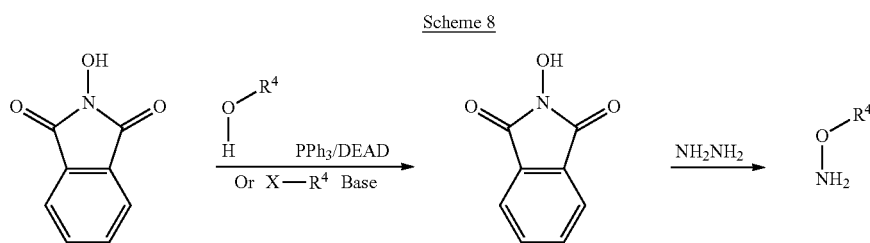
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carboxylate, ethyl 2-chloro-4-amino-5-carboxylate, ethyl 2,4-dichloropyrimidine-5-carboxylate) or pyrimidyl-6 (ethyl 2-chloro-4-amino-pyrimidine-6-carboxylate, ethyl 4,5-diaminopyrimidine 6-carboxylate) which are also considered as examples of 6-membered heterocyclic rings as R6 (Tetrahedron Lett. 1967, 1099; Chem Pharm. Bull. 1970, 1003. Justus Liebig Ann. Chem. 1954, 45).

B-3) The hydroxylamines required in both above schemes 6 and 7, can be prepared in several methods, as outlined in scheme 8 below.

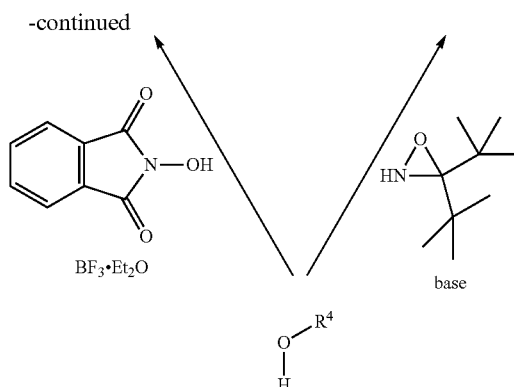
These hydroxylamines may be prepared firstly according to J. Antibiotics, 2000, 1072 with N-hydroxyphthalimide via Mitsunobu reaction conditions in presence of the alcohols R4-OH or by alkylation of N-hydroxyphthalimide in presence of activated compounds R4-X (X can be halogens like Cl, Br, I, or activated sulfonate esters like mesylate, tosylate, triflate, etc.). When R4 has the structure $\text{C}(\text{R}_x)(\text{R}_y)/\text{Z}$, with both R_x and R_y different from hydrogen, or is a tertiary alkyl, then N-hydroxyphthalimide may simply be treated with a stoichiometric amount of $\text{BF}_3 \cdot \text{Et}_2\text{O}$ with the corresponding alcohol R4-OH (Tetrahedron Lett., 2005, 6667). Formation of the final hydroxylamines may be performed in presence of

either hydrazine or methyl hydrazine. The oxaziridine technology developed by Ellman can also been employed to give directly deprotected O-substituted hydroxylamines (J. Org. Chem. 1999, 6528). (scheme 8).



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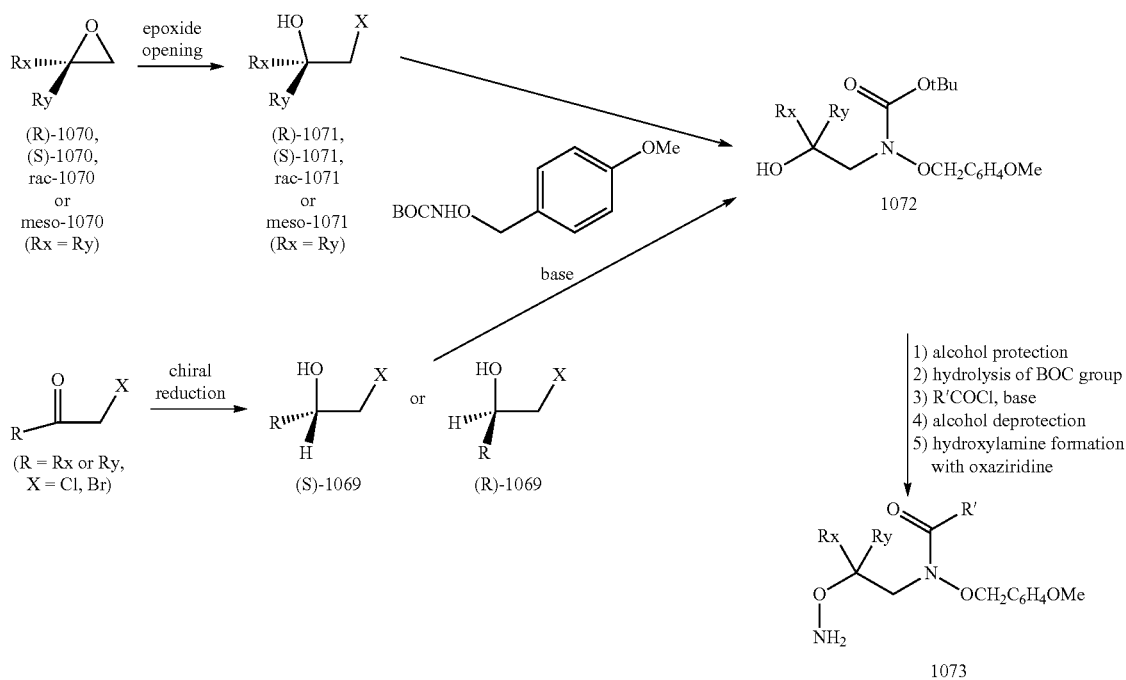
Alcohols R4-OH or compounds R4-X are either commercially available or may be prepared as described in following sections B-3-1a) to B-3-1e).

B-3-1a) A first method, where R4 is C(Rx)(Ry)Z, with Z=CH₂N(OH)COR', is outlined, as example, in the following scheme B-3-1a, using α-halogenoalcohols 1069 or 1071 and N-Boc-O-(p-methoxybenzyl)hydroxylamine. The α-halogenoalcohols 1071 can be prepared from derivatized epoxides 1070 which can be opened in acidic conditions (Eur. J. Org. Chem. 2004, 2557). The α-halogenoalcohols 1069 can be obtained by chiral reduction of corresponding α-halogenoketones (Tetrahedron:Asymmetry 2005, 3955).

epoxide forming reaction such as the catalytic epoxidation developed by Aggarwal (Accounts of Chemical Research, 2004, 37, pp. 611ff.). The conversion of 1069 and 1071 to 1072 is analogous to the procedure of Bioorganic & Medicinal Chemistry Letters 1996, 6(17), 2077ff. In the conversion of 1072 to 1073, again the oxaziridine technology developed by Ellman can be employed to give the hydroxylamine (J. Org. Chem. 1999, 6528).

B-3-1b) A second method, particular suited for hydroxylamines where R4 is C(Rx)(Ry)COOH, is outlined in the following scheme B-3-1b. These hydroxylamines may be obtained from the corresponding alcohols R4-OH via Mit-

Scheme B-3-1a

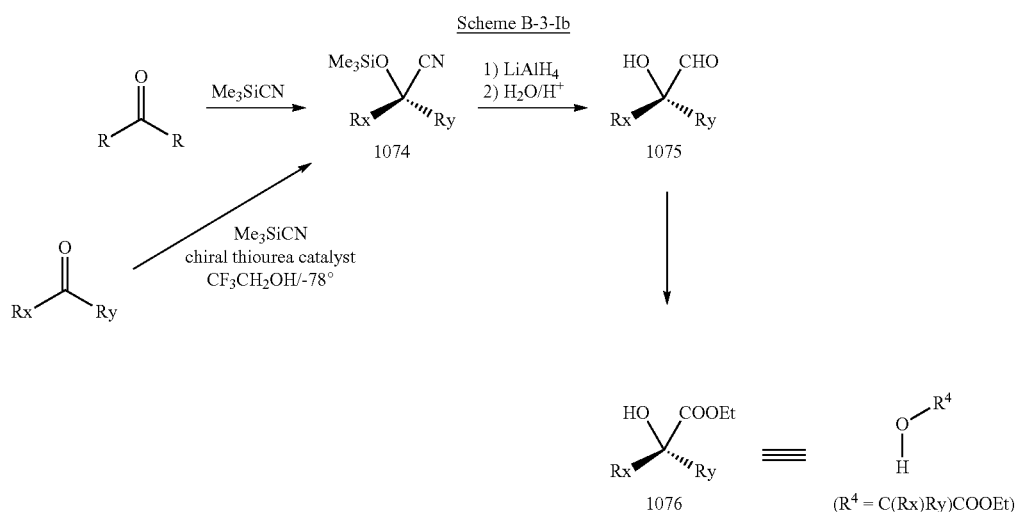


In the above scheme B-3-1a, the epoxides meso-1070 may be made from a ketone with two identical residues Rx=Ry=R with trimethylsulfonium iodide and base. For the epoxides (R)-1070 and (S)-1070 the synthesis may start from a ketone with two different residues Rx≠Ry, using an asymmetric

sunobu reaction conditions in presence of N-hydroxyphthalimide. The starting material R4-OH in turn may be obtained, as shown in the upper parts of scheme B-3-1b, from an appropriately Rx,Ry-substituted ketone. To this ketone is added trimethylsilyl cyanide, either with or without a chiral catalyst,

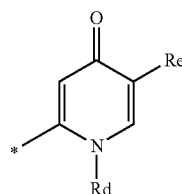
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to obtain a silyl protected cyanohydrin 1074. The chiral thio-urea catalyst indicated in scheme B-3-Ib, if used, and its conditions for use have been described in J. Am. Chem. Soc. 2005, 8964. The protected cyanohydrin is then reduced with LiAlH_4 to the corresponding aldehyde 1075 and then esterified to 1076. As an alternative to the Mitsunobu reaction using R4-OH, the hydroxyl group of N-hydroxyphthalimide can also be alkylated in presence of activated compounds R4-X (X can be halogens like Cl, Br, I, or activated sulfonate esters like mesylate, tosylate, triflate, etc; these are easily obtainable from the corresponding R4-OH). Formation of the desired hydroxylamine may be performed in presence of either hydrazine or methyl hydrazine, as shown in the lower part of scheme B-3-Ib, following the procedure described in J. Antibiotics, 2000, 1071.



B-3-Ic) A further variant for the preparation of hydroxylamines where R4 is $\text{C}(\text{Rx})(\text{Ry})\text{COOH}$, may start with the epoxides 1070 shown in above scheme B-3-Ia. These epoxides may be opened, as is customary, with aqueous base to form a vicinal diol; in which the primary hydroxy group is then converted to the aldehyde and then to the ester as outlined in the above scheme B-3-Ib in the conversion of 1075 to 1076.

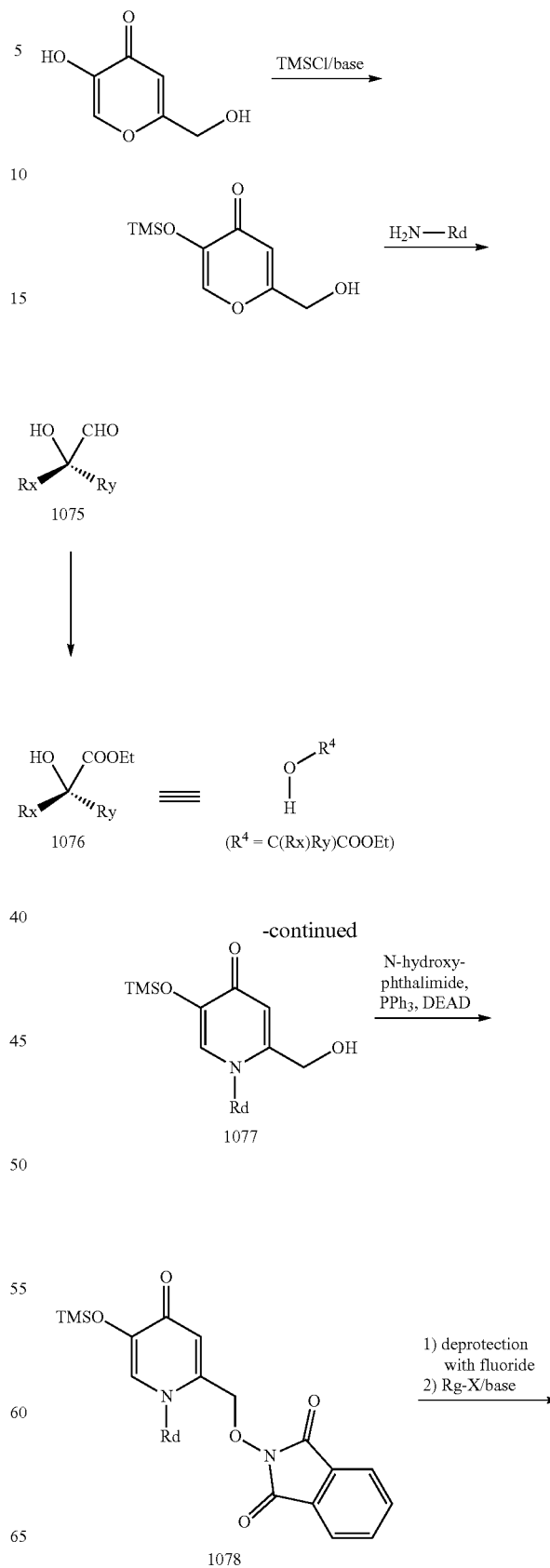
B-3-Id) For hydroxylamines where in R4 Z is

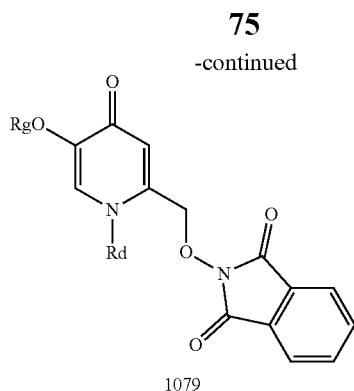


wherein Re and Rd are ORg (Rg being preferably selected from hydrogen, alkyl or optionally substituted benzyl for each Re and Rd), Rd is preferably hydrogen, alkyl, amino, monoalkylamino, optionally substituted benzyl, alkoxy-carbonyl or ORg (wherein Rg is as for Re and Rd); and wherein $\text{Rx}=\text{Ry}=\text{H}$; the following synthetic scheme B-3-Id may be adopted:

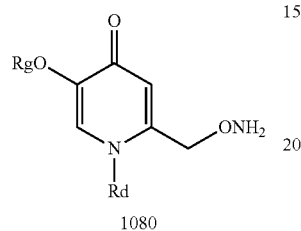
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Scheme B-3-Id



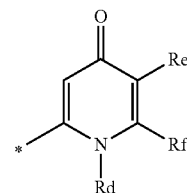


hydrazone

**76**

amines, when Rd is hydrogen, alkyl or optionally substituted benzyl, they are hydrazines, when Rd is amino or monoalkylamino; they are carbamates, when Rd is alkoxycarbonyl; and when Rd is ORg, these reactands are hydroxylamines. In the conversion of 1078 to 1079, the reactands Rg—X are hydrogen halides, alkyl halides or benzyl halides, wherein X is preferably Br or I. These reactands Rg—X are known or easily made from the corresponding alcohols Rg—OH. In the above scheme, instead of trimethylsilyl other protecting groups could also be used, such as benzyl, diphenyl or trityl as reported in J. Antibiotics 1990, 1450.

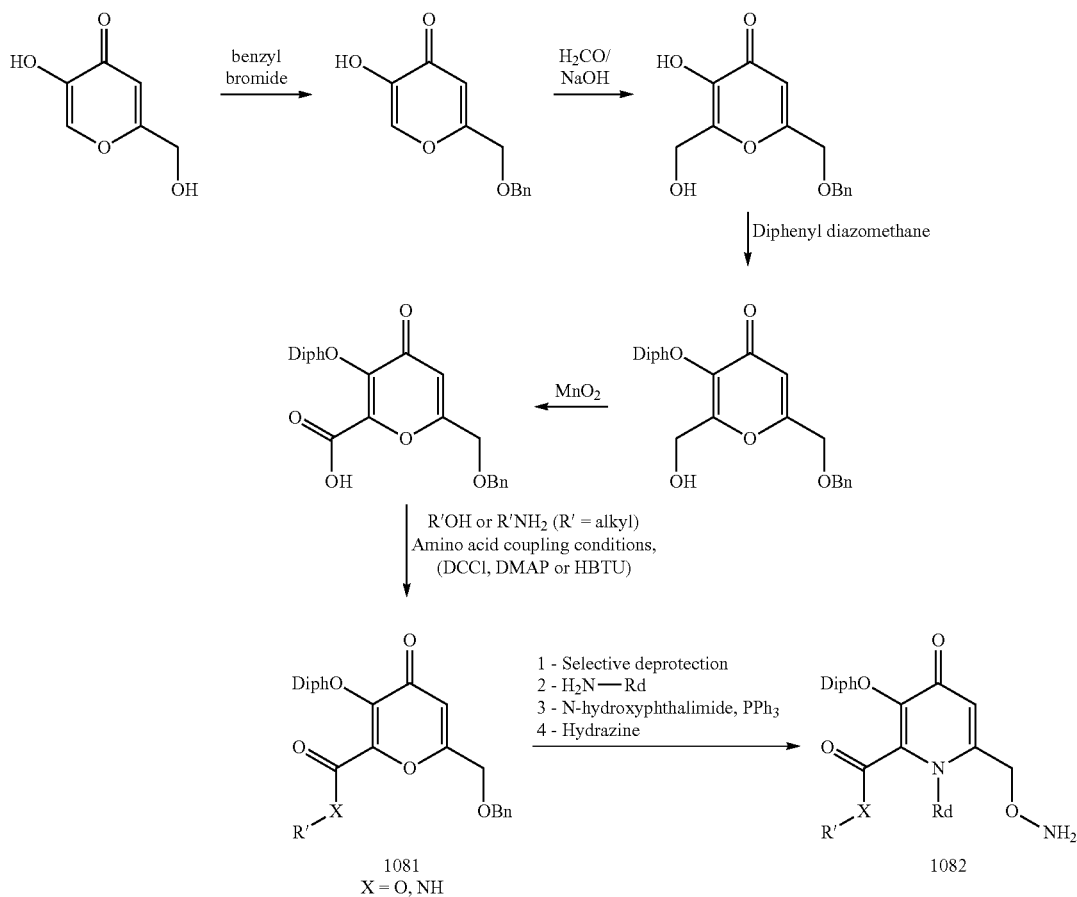
B-3-Ie) For hydroxylamines where in R4 Z is



This scheme adheres to the synthetic procedure described in EP-A-0 251 299. The kojic acid used as the starting material is commercially available. In the conversion from the silyl-protected kojic acid to 1077 the reactands H₂N-Rd are

wherein Re and Rd are ORg (Rg being preferably selected from hydrogen, alkyl or optionally substituted benzyl for each Re and Rd), Rf is preferably alkoxycarbonyl or alkylaminocarbonyl; and wherein Rx=Ry=H; the following synthetic scheme B-3-Ie may be adopted:

Scheme B-3-Ie

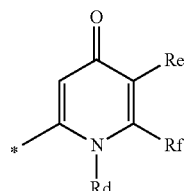


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The synthesis starts from kojic acid and is based on the known chemistry described in Biorg Med Chem Lett 2004, 3257; J. Med Chem 2002, 633; Bioorg Med. Chem. 2001 563 and J. Antibiotics 1990 1454. In the above scheme, the diphenylmethane protecting group in 1082 may subsequently be removed and, if desired, the free hydroxy group be reacted with an appropriate halide R—X, wherein R is alkyl or optionally substituted benzyl. These last steps are not shown in the above scheme.

B-3-If) For hydroxylamines where in R4 Z is

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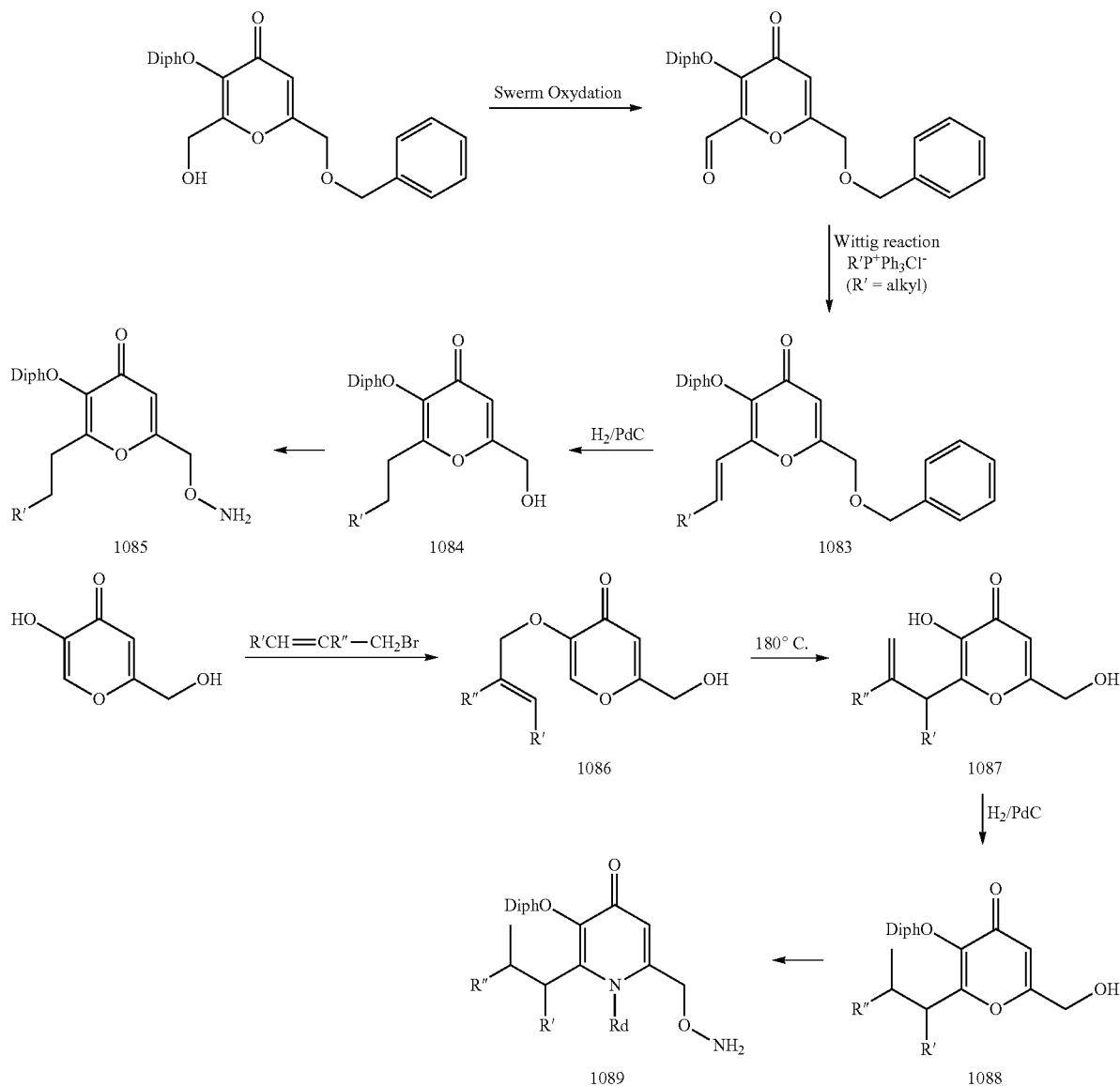


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wherein Re and Rd are ORg (Rg being preferably selected from hydrogen, alkyl or optionally substituted benzyl for each Re and Rd), Rf is preferably alkyl; and wherein Rx=Ry=H; a gain starting from kojic acid and with adapted protecting group strategy, further examples of pyridone derivatives which can be prepared are according to following scheme B-3-If:

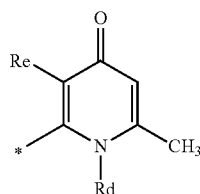
Scheme B-3-If



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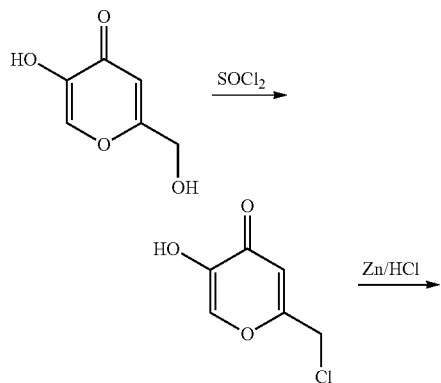
The upper part of scheme B-3-Ig is based on the possibility to run a Wittig reaction (J. Med. Chem 2004, 6349) on the aldehyde obtained after Swern oxidation. The resulting product 1083 may be subjected to hydrogenolysis in presence of Pd/C similarly to the preparation of compound 10'j in J. Med. Chem 2004, 6349. The resulting compound 1084 may then be treated as already outlined in above scheme 8 to obtain desired hydroxylamines 1085. The lower part of scheme B-3-Ig shows the preparation of hydroxylamines with linear or branched alkyl as Rf. In this part of scheme, R' and R'' are preferably selected from alkyl (in particular methyl and ethyl) and hydrogen; more preferably one of R and R'' is hydrogen, or both R' and R'' are hydrogen. A thermal rearrangement of products 1086, obtainable by alkylation of the 4-hydroxyl group of kojic acid with appropriately R',R''-substituted allyl bromides (J. Am. Chem. Soc. 1956, 2816) leads to 6-substituted pyridones 1087. These may again be converted in the usual way to hydroxylamines 1089. In the above formed hydroxylamines 1085 and 1089 the diphenylmethane group may subsequently be removed and, if desired, the free hydroxy group be reacted with an appropriate halide R—X, wherein R is alkyl or optionally substituted benzyl. These last steps are not shown in the above scheme.

B-3-Ig) For hydroxylamines where in R4 Z is



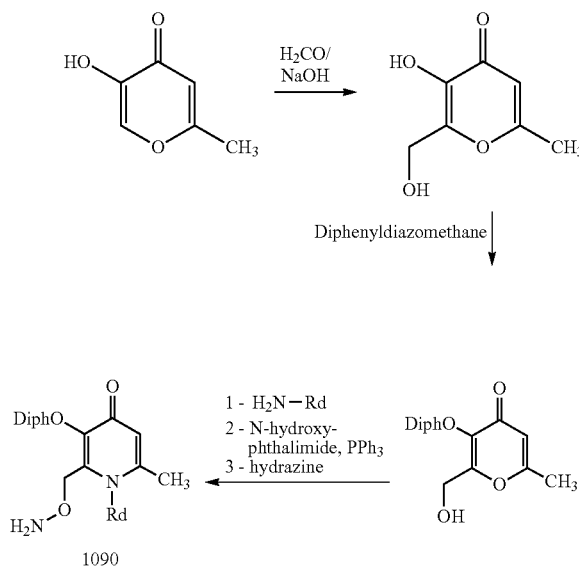
wherein Re is ORg (Rg being preferably selected from hydrogen, alkyl or optionally substituted benzyl), and Rd is preferably selected from hydrogen, alkyl, amino, monoalkylamino, optionally substituted benzyl or alkoxy carbonyl; and wherein Rx=Ry=H; a synthesis analogous to the scheme B-3-Ig shown below may be used. Details of the syntheses in the upper part of this scheme can be found in J Med. Chem. 2004, 6349.

Scheme B-3-Ig



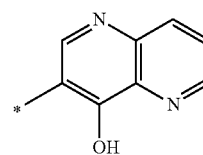
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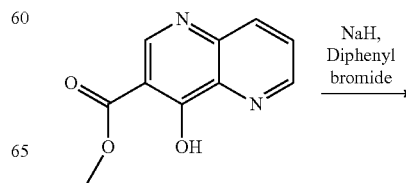
In the hydroxylamines 1090 obtained in above scheme B-3-Ig the diphenylmethane group may subsequently be removed and, if desired, the free hydroxy group be reacted with an appropriate halide R—X, wherein R is alkyl or optionally substituted benzyl. These last steps are not shown in the above scheme.

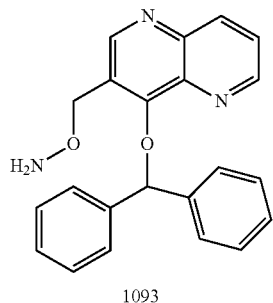
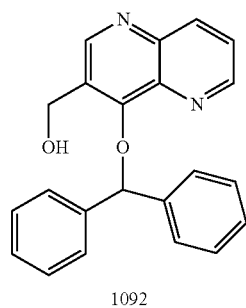
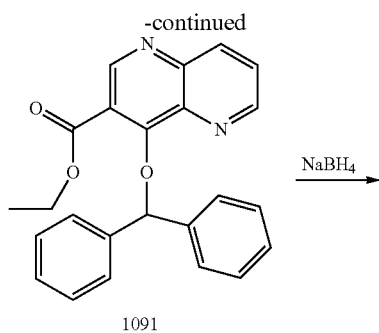
B-3-II) For hydroxylamines where Z is



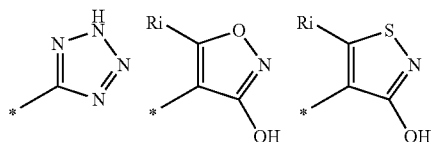
the following scheme B-3-II may be used. In this scheme, the naphthyridine hydroxylamine may be obtained from commercially available 4-hydroxy-1,5-naphthyridine-3-carboxylic ethyl ester after diphenyl protection of the hydroxyl group and reduction of ester group. Mitsunobu reaction in presence of N-hydroxyphthalimide and deprotection with hydrazine leads to the desired compound 1093.

Scheme B-3-II



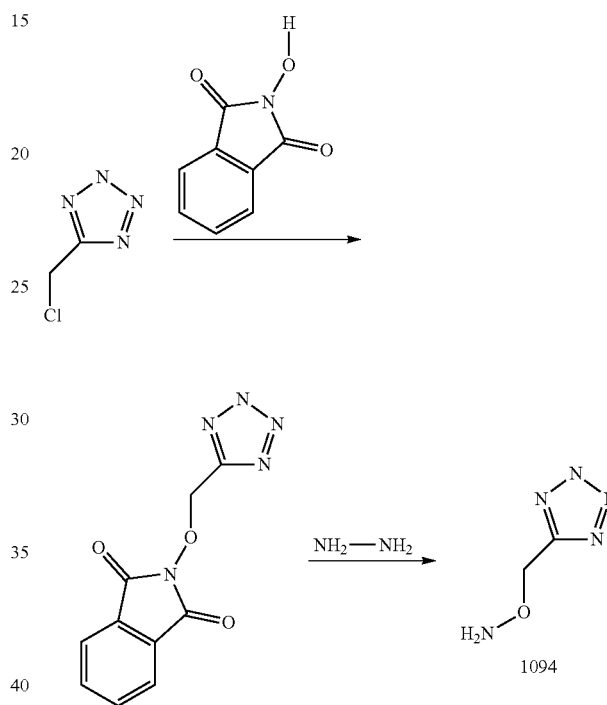
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B-3-III) Further suited hydroxylamines are those where Z is selected from:

**82**

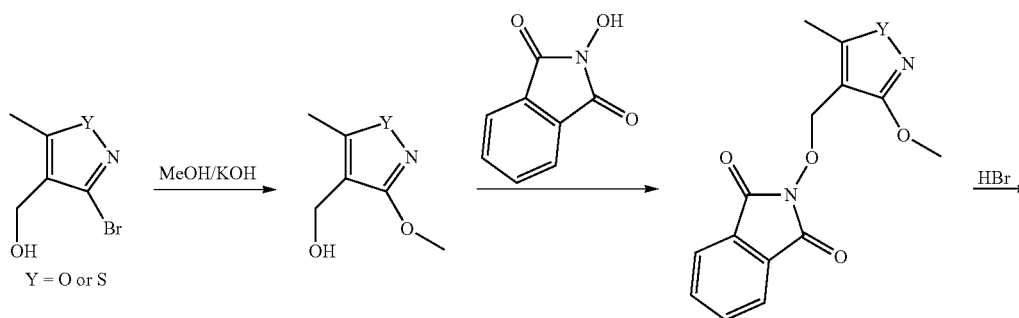
wherein Ri is as defined in claim 1. Those substituted 5-membered rings are known as bioisosteres (Curr. Med. Chem. 2004, 11, 945) of the carboxylic acid group. One particularly preferred hydroxylamine of this type, tetrazole methyl hydroxylamine 1094, may be obtained from commercially chloro methyl tetrazole with N-hydroxyphthalimide according to the method described in J. Antibiotics 2000, 1071 (scheme B-3-IIIa below).

Scheme B-3-IIIa



Isoxazole analogs (J. Heterocyclic Chem. 1997, 345, J. Med. Chem. 1996, 183) or isothiazoles (J. Med. Chem. 1998, 930) are also known as bioisosteres of the carboxylic acid group and may be done in similar way as outlined in the scheme B-3-IIIb below based on the following literature (J. Chem. Soc., Perkin 1 1993, 2153; Synthesis 1996, 1177, Acta Chem. Scand. 1990, 96).

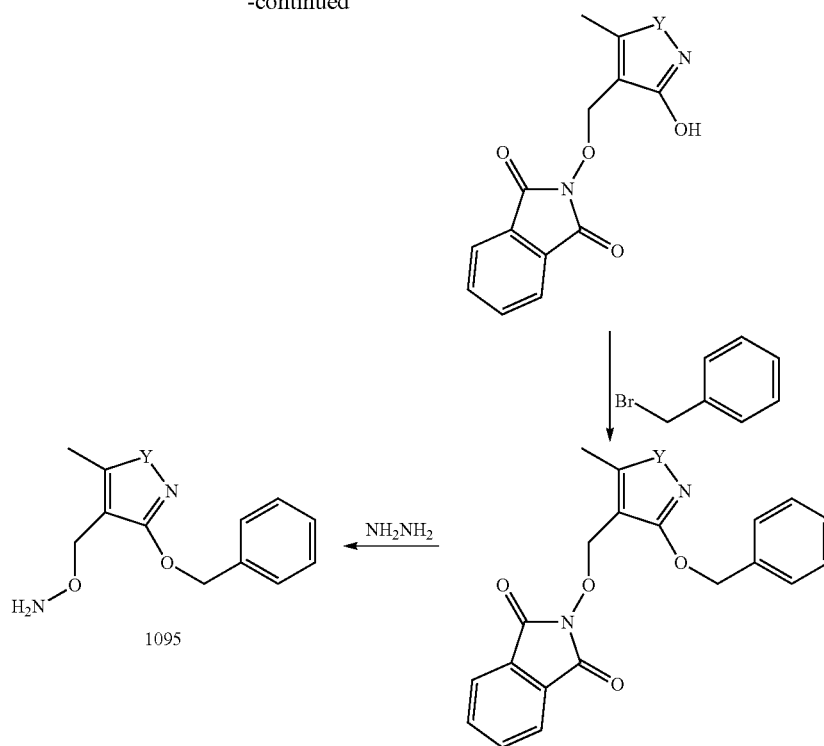
Scheme-B-3-IIIb



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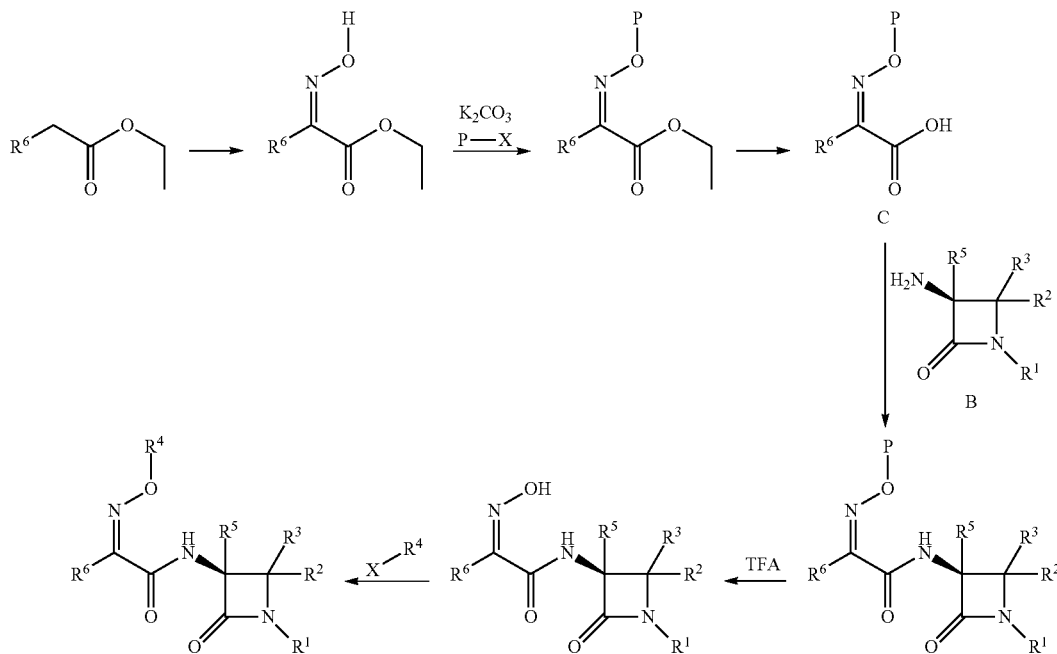
In this scheme, the removal of the benzyl protecting group in the resulting hydroxylamine-containing isoxazoles and isothiazoles 1095 is not shown.

After formation of the ketoacid derivative A3, the condensation with O-substituted hydroxylamine (prepared or commercially available like O-methyl hydroxylamine) may be performed to lead to compounds of general formula A. The condensation of the hydroxylamines R⁴-ONH₂ with the keto acid derivatives A3 to form the compounds A may then follow

the procedure described in J. Antibiotics 2000, 1071 and WO-A-02/22613.

B-4) As an alternative of preparing compounds A, the esters A1 may firstly be oxidised to form compounds A2 as shown in scheme 6, then reacted with hydroxylamine, followed by O-protection and ester hydrolysis, to lead to the intermediate C, which can be coupled to the azetidinone B. Substituents R₄ can be introduced by alkylation. This alternate route is shown in the following scheme 9:

Scheme 9



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The compounds of formula II are compounds known from the above cited literature references (EP-A-0 508 284 and U.S. Pat. No. 6,566,355) or can be made in an analogous manner or can be made as described in the following scheme 10.

The intermediate compound D (J. Med. Chem. 1998, 3961 and EP-A-0 508 234) gives access to compounds of formula IIA, IIB or to IIC using different synthetic routes outlined in scheme 10.

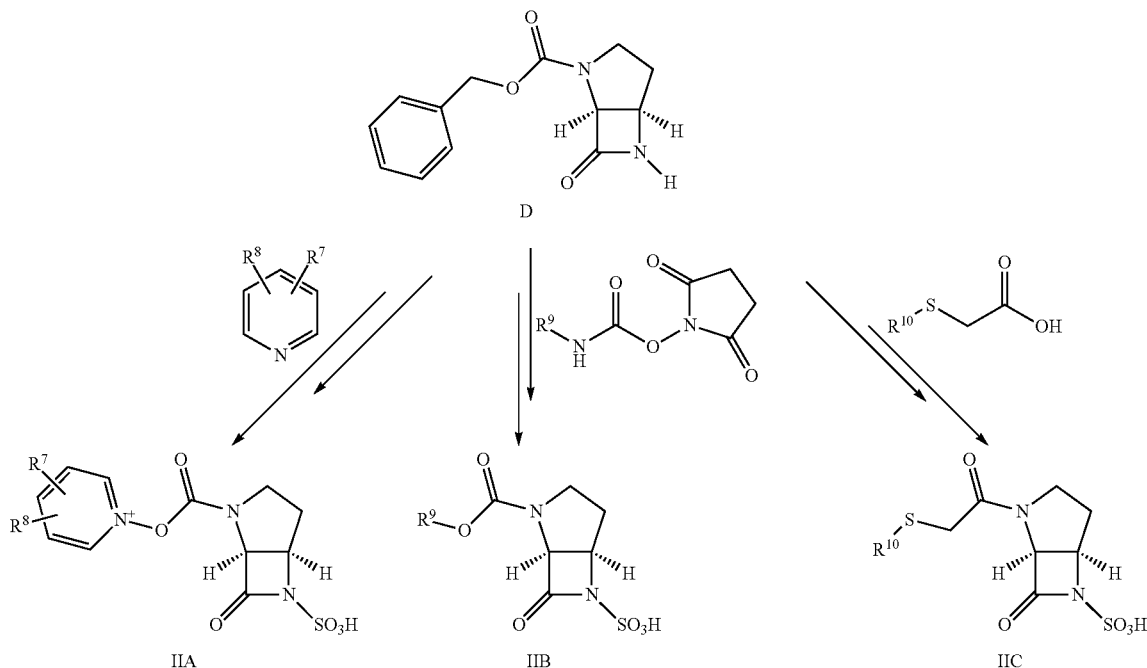
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1982, 5160). Pyridine derivatives, condensed at room temperature in dimethylformamide, are either commercially available or synthesized according to known literature procedures.

Second pathway: Compound D may be first hydrogenated in presence of BOC_2O to afford the intermediate G (Tetrahedron Lett. 1988, 2983). Then sulfonation of compound G followed by removal of the BOC protect-

Scheme 10



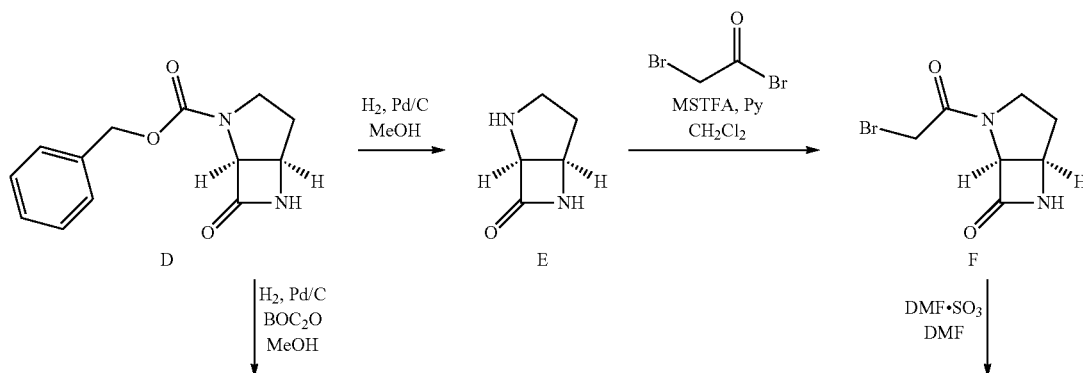
For the preparation of pyridinium carboxymethyl derivatives of formula IIA, two synthetic routes are possible (scheme 11):

First pathway: Compound F, prepared from compound D in presence of bromo acetyl bromide according to the procedures described in J. Med. Chem. 1998, 3961 and EP-A-0 508 234, may be sulfonated (J. Org. Chem.

ing group generates compound H (J. Med. Chem. 1998, 3961 and J. Org. Chem. 1982, 5160).

At that stage the pyridinium carboxymethyl derivatives (prepared by analogy according to the procedures described in Synthesis 2000, 1733 or J. Chem. Soc., Perkin Trans. 11977, 1692) may be introduced to generate compounds IIA.

Scheme 11

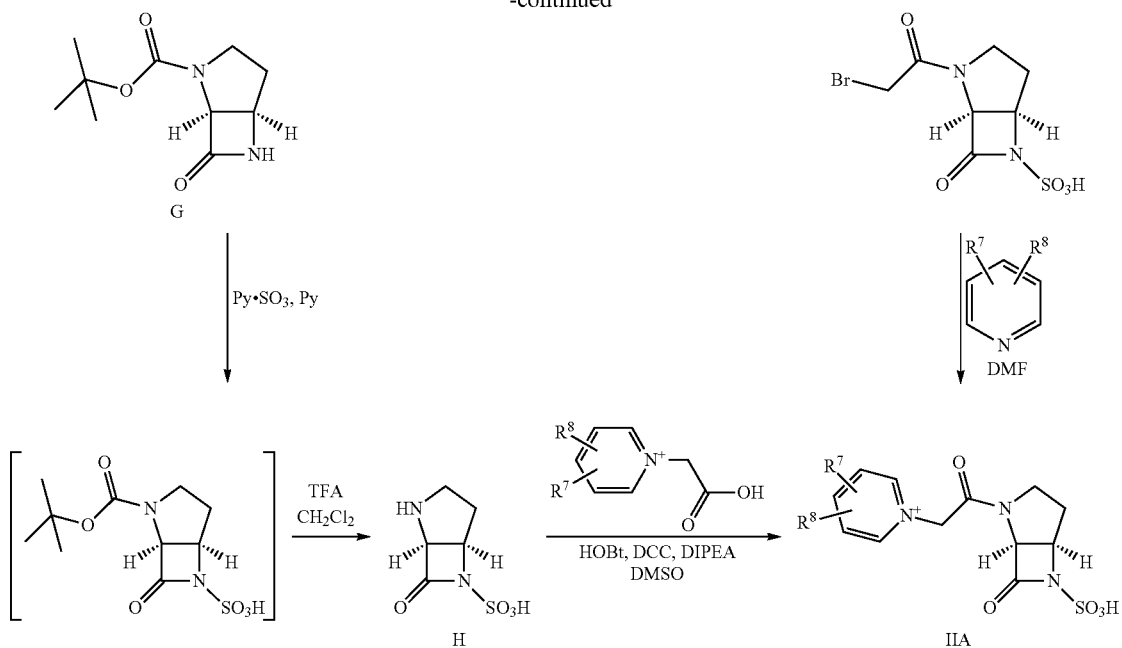


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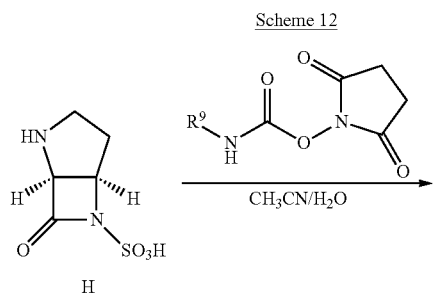
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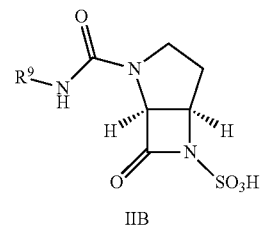


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The compounds of formula IIB may be obtained from compound H and succinimidyl derivatives according to the following scheme 12:

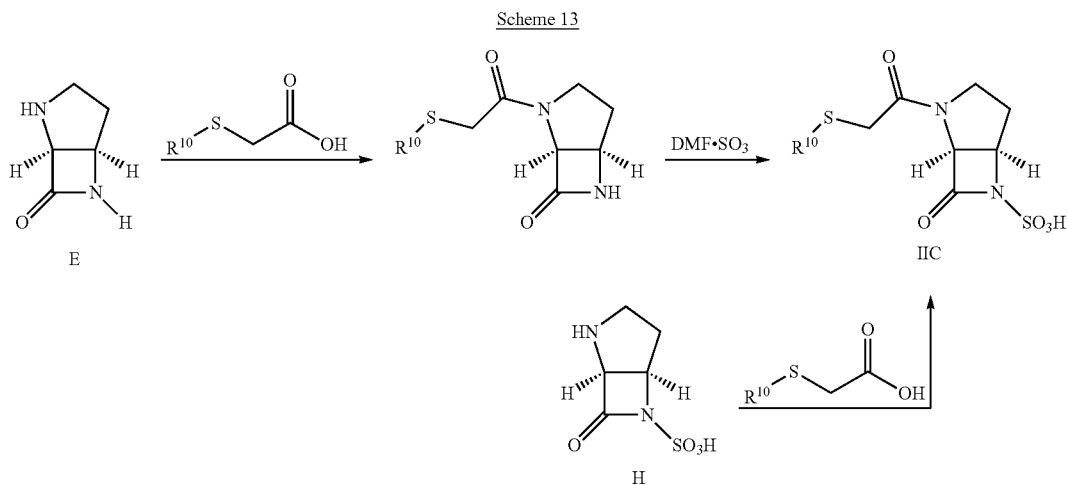


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Compound H may be prepared as previously described in scheme 11. Then the succinimidyl derivatives may be synthesized and introduced according to the procedures described in J. Med. Chem. 1998, 3961.

Compounds of formula IIC may be obtained either from compound E or compound H according to the following scheme 13:



The compounds IIC may be synthesized via two different routes:

- either from compound E by first coupling with the thioacetic acid derivatives, followed by a sulfonation step (J. Med. Chem. 1998, 3961),
- or directly from compound H (see scheme 11) and thioacetic acid derivatives.

The preparation of sodium salt of compound of formula I and II can be performed either with the procedures described in WO-A-02/22613, U.S. Pat. No. 6,566,355, J. Med. Chem. 1998, 3961 or in J. Antibiotics, 1985, 346.

In the above descriptions, the reactants are reacted together with a suitable solvent at elevated or low temperatures for sufficient time to allow the reaction to proceed to completion. The reaction conditions will depend upon the nature and reactivity of the reactants. Wherever a base is used in a reaction, they are selected from e.g. triethylamine, tributylamine, trioctylamine, pyridine, 4-dimethylaminopyridine, diisopropylamine, 1,5-diazabicyclo[4,3,0]non-5-ene, 1,8-diazabicyclo[5,4,0]undec-7-ene, sodium carbonate, sodium dicarbonate, potassium carbonate, potassium bicarbonate or cesium carbonate.

The deprotection of functional groups may be carried out either by hydrogenation or hydrolysis with appropriate acids such as hydrochloric acid, formic acid, trifluoroacetic acid, acetic acid or p-toluenesulfonic acid; in solvents such as methanol, ethanol, propanol, ethyl acetate, acetonitrile, methylene chloride or ethylene chloride. The hydrogenation is usually carried out in the presence of a metal catalyst, such as Pd, Pt or Rh under normal to high pressure.

The solvents of choice for the reaction are selected based upon the reactants used and from solvents such as benzene, toluene, acetonitrile, tetrahydrofuran, ethanol, methanol, chloroform, ethyl acetate, methylene chloride, dimethyl formamide, dimethyl sulfoxide, hexamethyl phosphoric triamide or the like. Solvents mixtures may also be used.

Reaction temperatures would generally range from between -70°C . to 150°C . The preferred molar ratio of the reactants is 1:1 to 1:5. The reaction time range from 0.5 to 72 hours, depending on the reactants.

The compounds of formula I, Ia and Ib and their pharmaceutically compatible salts can be used in accordance with the invention in the control or prevention of illness in mammals, human and non-human, especially in combination with β -lactamase inhibitors.

Thereby, the compound of formula I or pharmaceutically compatible salts thereof with bases can be administered before, simultaneously with or after the administration or intake of one or more β -lactamase inhibitors of formula II-XIII. The products in accordance with the invention can be administered in the form of pharmaceutical compositions containing the combination of a compound of formula I or a pharmaceutically compatible salt thereof with a base, and one or more β -lactamase inhibitors of formula II-XIII; alternatively, they may also be administered separately from the β -lactamase inhibitors, simultaneously or sequentially, in which case the combination according to the invention may be present as a kit-of-parts. Articles with such pharmaceutical combinations are also an object of the present invention.

The compounds of formula I are active against a variety of bacterial organisms. They are active against aerobic Gram-negative bacteria that do not produce β -lactamases, including Enterobacteriaceae, for example *Escherichia coli*, *Enterobacter cloacae*, *Enterobacter aerogenes*, *Citrobacter freundii*, *Klebsiella pneumoniae*, *Klebsiella oxytoca*, *Proteus vulgaris*, *Providencia rettgeri*; *Pseudomonas* for example *P. aeruginosa*; *Acinetobacter* for example *A. baumannii*;

Burkholderia, for example *B. cepacea*; *Stenotrophomonas* for example *S. maltophilia*. Combinations of compounds of formula I and formula II are active against strains of the above organisms that do produce β -lactamases and this activity can be increased by additionally combining compounds of formula III-XIII with the combination comprising compounds of formulae I and II.

Formulations

The pharmaceutical compositions and articles (kits-of-parts) according to the present invention are administered by any route, preferably in the form of a pharmaceutical composition, or kit-of-parts of individual compositions, adapted to such a route. Dosage and route of administration should be determined by susceptibility of the causative organisms, severity and site of infection, and the condition of the patient. The preferred types of pharmaceutical compositions are, for example, administered intravenously or by intramuscular injection.

Formulations for parenteral administration can be in the form of aqueous isotonic sterile injection solutions or suspensions. These solutions or suspensions can be prepared from sterile powders, granules or lyophilizates. The compounds can be dissolved in sterile water or in various sterile buffers that may contain sodium chloride, polyethylene glycol, propylene glycol, ethanol, sucrose, glucose, arginine, lysine, or lactic acid. The dry compositions can contain from 0.1% to 99% by weight, preferably 10%-60% by weight, of each of active ingredients. If the compositions contain dosage units, each unit preferably contains from 50 mg to 4 g of each active substance.

The ratio of β -lactam antibiotic (compounds of formula I or pharmaceutically compatible salts thereof with a base) and β -lactamase inhibitors (compounds of formula II and formula III-XIII, or pharmaceutically compatible salts thereof with a base) can also vary within wide limits and will be fitted to the individual requirements in each particular case. In general, a ratio of between 1 part of antibiotic of general formula I to 5 parts of any one β -lactamase inhibitor of general formula II or III-XIII and 20 parts of antibiotic of general formula I to 1 part of any one β -lactamase inhibitor of general formula II or III-XIII should be appropriate.

The dosage of the compound of formula I and of the pharmaceutically compatible salts thereof with bases can vary within wide limits and will be fitted in each particular case to the individual requirements and to the β -lactamase producing pathogen to be controlled. In general, a dosage of about 0.1 to about 2 g of antibiotic administered one to four times over a 24 hours period should be appropriate.

The present invention is further illustrated by the following non-limiting examples.

Example 1

(3S,4S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(5-hydroxy-1-methyl-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidine-sulfonic acid (6)

Preparation of 5-(diphenylmethoxy)-2-(hydroxymethyl)-1-methylhydropyridin-4-one

5-(Diphenylmethoxy)-2-(hydroxymethyl)pyran-4-one (J. Antibiotics 1990, 189) (5.0 g, 16.22 mmol) and methyl amine (80.6 g, 1.04 mol) were stirred at room temperature overnight in presence of methanol (1 mL). The precipitate observed was filtered off and the mother liquor was extracted 3 times with

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ethyl acetate. The organic phases were dried and the solvent evaporated. The total amount of collected product was 2.1 g.

¹H-NMR (DMSO-d₆) δ: 3.49 (s, 3H), 4.29 (s, 2H), 6.22 (s, 1H), 6.74 (s, 1H), 7.20-7.60 (m, 11H).

Preparation of 2-[[5-(diphenylmethoxy)-1-methyl-4-oxo-2-hydropyridyl]methoxy]benzo[c]azoline-1,3-dione

In THF (10.0 mL), containing triphenylphosphine (0.470 g, 1.79 mmol) and N-hydroxyphthalimide (0.293 g, 1.79 mmol) was added 5-(diphenylmethoxy)-2-(hydroxymethyl)-1-methylhydropyridin-4-one (0.240 g, 0.75 mmol). After cooling the solution at 0° C., diethyl-azodicarboxylate (0.312 g, 1.79 mmol) was added dropwise and stirred for 30 min at this temperature. The solution was then warmed up to room temperature and stirred overnight. A suspension was observed, filtered off and washed to give 210 mg of the desired compound.

¹H-NMR (DMSO-d₆) δ: 3.78 (s, 3H), 5.08 (s, 2H), 6.29 (s, 1H), 6.74 (s, 1H), 7.20-7.50 (m, 10H), 7.69 (s, 1H), 7.80-7.90 (m, 4H),

Preparation of 2-[(aminooxy)methyl]-5-(diphenylmethoxy)-1-methylhydropyridin-4-one

Hydrazine hydrate (0.023 mL, 0.47 mmol) was added to ethanol (10 mL) already containing 2-[[5-(diphenylmethoxy)-1-methyl-4-oxo-2-hydropyridyl]methoxy]benzo[c]azoline-1,3-dione (0.200 g, 0.43 mmol). The resulting solution was refluxed for 2 hours. After cooling at room temperature the precipitate was collected and the ethanol was evaporated. The resulting residue was triturated in ethyl acetate to give 130 mg of the expected compound.

¹H-NMR (DMSO-d₆) δ: 3.51 (s, 3H), 4.43 (s, 2H), 6.21 (s, 1H), 6.29 (br s, 2H), 6.74 (s, 1H), 7.20-7.50 (m, 10H), 7.56 (s, 1H).

Preparation of (2Z)-3-[[5-(diphenylmethoxy)-1-methyl-4-oxo(2-hydropyridyl)]methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid

In a mixture of ethanol (5 mL)/chloroform (3 mL), 2-[(aminooxy)methyl]-5-(diphenylmethoxy)-1-methylhydropyridin-4-one (0.076 g, 0.23 mmol) and 2-oxo-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}acetic acid (0.085 g, 0.21 mmol) were stirred at room temperature for 12 hours. The solvents were evaporated and ethyl acetate was added to the residue. The resulting suspension was filtered off to afford 77 mg of the desired compound.

¹H-NMR (DMSO-d₆) δ: 3.49 (s, 3H), 5.00 (s, 2H), 6.24 (s, 1H), 6.72 (s, 2H), 6.83 (s, 1H), 7.15-7.50 (m, 25H), 7.57 (s, 1H), 8.85 (s, 1H).

Preparation of (3S,4S)-3-((2Z)-3-[[5-(diphenylmethoxy)-1-methyl-4-oxo(2-hydropyridyl)]methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoylamino)-4-methyl-2-oxoazetidinesulfonic acid

(2Z)-3-[[5-(diphenylmethoxy)-1-methyl-4-oxo(2-hydropyridyl)]methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid (0.380 g, 0.52 mmol), dicyclohexylcarbodiimide (0.160 g, 0.78 mmol) and 1-hydroxy-7-azabenzotriazole (0.106 g, 0.78 mmol) were stirred at room temperature for 3 hours. Then (3S,4S)-3-amino-4-

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methyl-2-oxoazetidinesulfonic acid (0.103 g, 1.04 mmol) and a catalytic amount of triethylamine were added to the previous solution which was stirred for 16 hours at room temperature. The solvent was evaporated in vacuo and the residue was purified by column chromatography (silica gel, eluent; dichloromethane and methanol, 95/5, v/v). 100 mg of desired compound was obtained.

¹H-NMR (DMSO-d₆) δ: 1.35 (d, 3H, J=6.0 Hz), 3.55-3.60 (m, 1H) 3.83 (s, 3H), 4.34 (dd, 1H, J=2.5, 7.7 Hz), 5.21 (s, 2H), 6.75 (m, 1H), 6.77 (s, 1H), 7.00 (s br, 1H), 7.15-7.60 (m, 26H), 8.39 (s br, 1H) 8.83 (s, 1H), 9.33 (d, 1H, J=7.7 Hz).

(3S,4S)-3-((2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(5-hydroxy-1-methyl-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino)-4-methyl-2-oxoazetidine-sulfonic acid (6)

(3S,4S)-3-((2Z)-3-[[5-(diphenylmethoxy)-1-methyl-4-oxo(2-hydropyridyl)]methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoylamino)-4-methyl-2-oxoazetidinesulfonic acid (0.076 g, 0.08 mmol) was dissolved in dichloromethane (3 mL). Triethylsilane (0.021 mL, 0.25 mmol) was added at -10° C. and trifluoroacetic acid (0.327 mL, 4.25 mmol) was added dropwise and stirred for 1 h at the same temperature. The solution reacted at room temperature for 2 hours. Then dichloromethane was evaporated in vacuo and the residue was purified by preparative HPLC to give 35 mg of the desired compound.

Example 2

(3S,4S)-3-((2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino)-4-methyl-2-oxoazetidinyl hydroxysulfonate (12)

The titled compound was prepared following scheme 1. (2Z)-3-[[[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl)]methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1990, 1450 and WO-A-02/22613 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinyl hydroxysulfonate was prepared according to the procedures described in J. Am. Chem. Soc, 1982, 6053 and J. Antibiotics, 1985, 1536.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 3

(3S,4S)-3-((2Z)-2-(4-aminopyrimidin-2-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino)-4-methyl-2-oxoazetidine-sulfonic acid (21)

The titled compound was prepared following scheme 8. (2Z)-3-[[[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl)]methoxy]-2-(4-[(triphenylmethyl)amino]pyrimidin-2-yl))-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1984, 546 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinesulfonic acid was prepared according to the procedures described in J. Org. Chem. 1980, 410.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

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Example 4

2-((3S,4S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinyloxy)acetic acid (22)

The title compound was prepared following scheme 1. (2Z)-3-{[1,5-Bis(diphenylmethoxy)-4-oxo(2-hydropyridyl))methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1990, 1450 and WO-A-02/22613 and 2-((3S,4S)-3-amino-4-methyl-2-oxoazetidinyloxy)acetic acid was prepared according to the procedure described in J. Med. Chem. 1985, 1447 and J. Antibiotics, 1985, 813.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 5

(2S)-2-((3S,4S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinyloxy)propanoic acid (23)

The titled compound was prepared following scheme 1. (2Z)-3-{[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl))methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1990, 1450 and WO-A-02/22613 and (2S)-2-((3S,4S)-3-amino-4-methyl-2-oxoazetidinyloxy)propanoic acid was prepared according to the procedure described in J. Med. Chem. 1985, 1447 and J. Antibiotics, 1985, 813.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 6

(3S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4,4-dimethyl-2-oxoazetidinyloxyhydroxysulfonate (26)

The title compound was prepared following scheme 1. (2Z)-3-{[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl))methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1990, 43, 1450 and WO-A-02/22613 and (3S)-3-amino-4,4-dimethyl-2-oxoazetidinyloxyhydroxysulfonate was prepared according to the procedure described in J. Org. Chem. 2003, 177 and Tetrahedron Lett., 1986, 2789.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 7

(3S,4S)-3-{(2Z)-2-(2-amino-5-chloro(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinyloxyhydroxysulfonate (29)

The titled compound was prepared following scheme 1. (2Z)-3-{[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl))methoxy]-2-{5-chloro-2-[(triphenylmethyl)amino](1,3-

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thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in J. Antibiotics, 1990, 1450 and WO-A-02/22613 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinyloxyhydroxysulfonate was prepared according to the procedure described in J. Am. Chem. Soc., 1982, 6053 and J. Antibiotics, 1985, 1536.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 8

(3S,4S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(5-hydroxy-1-methoxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinesulfonic acid (5)

The title compound was prepared following scheme 1.

The alcohol derivative 5-(diphenylmethoxy)-2-(hydroxymethyl)-1-methoxyhydropyridin-4-one was obtained according to the following procedure;

5-(diphenylmethoxy)-2-(hydroxymethyl)-1-methoxyhydropyridin-4-one

To a solution of DMF (20 mL) containing 5-(diphenylmethoxy)-1-hydroxy-2-(hydroxymethyl)hydropyridin-4-one (J. Antibiotics 1990, 1450) (2.0 g, 6.19 mmol) at 0° C. was first added potassium tert-butoxide (0.971 g, 8.66 mmol) and then iodo methane (4.23 g, 8.66 mol). The resulting mixture was stirred for 30 min at 0° C. and then 2 h at room temperature. Then ethyl acetate (20 mL) and water (50 mL) was added. The observed precipitate was filtered off and the washed with additional ethyl acetate. The total of amount of collected product was 1.4 g.

¹H-NMR (DMSO-d₆) δ: 3.87 (s, 3H), 4.38 (d, 2H, J=5.8 Hz), 5.55 (t, 1H, J=5.8 Hz) 6.13 (s, 1H), 6.69 (s, 1H), 7.20-7.50 (m, 10H), 7.88 (s, 1H).

The compound of formula A (2Z)-3-{[5-(diphenylmethoxy)-1-methoxy-4-oxo(2-hydropyridyl))methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in example 1 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinesulfonic acid was prepared according to the procedures described in J. Org. Chem. 1980, 410.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 9

(3S,4S)-3-{(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[(1-hydroxy-5-methoxy-4-oxo(2-hydropyridyl))methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinesulfonic acid (7)

The titled compound was prepared following scheme 1.

2-(hydroxymethyl)-5-methoxypyran-4-one was prepared from kojic acid according to the procedure described in J. Org. Chem. 1950, 221

Then the preparation of the compound of formula A (2Z)-3-{[1-(diphenylmethoxy)-5-methoxy-4-oxo(2-hydropyridyl))methoxy]-2-{2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)}-3-azaprop-2-enoic acid was prepared according to the procedures described in example 1 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinesulfonic acid was prepared according to the procedures described in J. Org. Chem. 1980, 410.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 10

(3S,4S)-3-[(2Z)-2-(2-amino(1,3-thiazol-4-yl))-3-[2-(hydroxyacetyl amino)ethoxy]-3-azaprop-2-enoylamino]-4-methyl-2-oxoazetidinesulfonic acid (10)

The titled compound was prepared following scheme 1.

The preparation of the compound of formula A (2Z)-3-[(N-[(4-methoxyphenyl)methoxy]acetyl amino)ethoxy]-2-[(triphenylmethyl)amino](1,3-thiazol-4-yl))-3-azaprop-2-enoic acid was prepared according to the procedures described in Bioorg. Med. Chem. Lett. 1996, 2077 and (3S, 4S)-3-amino-4-methyl-2-oxoazetidinesulfonic acid was prepared according to the procedures described in J. Org. Chem. 1980, 410.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 11

(3S,4S)-3-{2-(2-amino(1,3-thiazol-4-yl))-3-[(1-amino-5-hydroxy-4-oxo(2-hydropyridyl)methoxy]-3-azaprop-2-enoylamino}-4-methyl-2-oxoazetidinesulfonic acid (11)

The titled compound was prepared following scheme 1.

1-[(1E)-2-(4-nitrophenyl)-1-azavinyl]-5-(diphenylmethoxy)-2-(hydroxymethyl)hydropyridin-4-one was prepared from kojic acid according to the procedure described in Helv. Chim. Acta 1960, 461

Compound of formula A 3-[(1-[(1E)-2-(4-nitrophenyl)-1-azavinyl]-5-(diphenylmethoxy)-4-oxo(2-hydropyridyl)methoxy)(2Z)-2-[(triphenylmethyl)amino](1,3-thiazol-4-yl))-3-azaprop-2-enoic acid was prepared according to the procedures described in example 1 and (3S,4S)-3-amino-4-methyl-2-oxoazetidinesulfonic acid was prepared according to the procedures described in J. Org. Chem. 1980, 410.

Final assembly and deprotection steps were done similarly according to the method described for the example 1.

Example 12

2-((3S,4S)-3-[(2Z)-2-(2-amino-5-chloro(1,3-thiazol-4-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl)methoxy]-3-azaprop-2-enoylamino)-4-[(aminocarbonyloxy)methyl]-2-oxoazetidinyloxy)acetic acid (42)

The titled compound was prepared following scheme 1.

The preparation of the compound of formula A (2Z)-3-[[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl)methoxy]-2-[5-chloro-2-[(triphenylmethyl)amino](1,3-thiazol-4-yl))-3-azaprop-2-enoic acid was achieved according to the procedures described in J. Antibiotics, 1990, 1450 and WO-A-02/22613 using 2-[(5-Chloro-2-[(triphenylmethyl)amino](1,3-thiazol-4-yl))-2-oxoacetic acid (obtained from 2-oxo-2-[(triphenylmethyl)amino](1,3-thiazol-4-yl)]acetic acid (DE2710902A1) and chlorination step achieved according to the procedures described in EP-A-0 055 465) and 2-[(aminooxy)methyl]-5-(diphenylmethoxy)-1-methylhydropyridine-4-one (described in example 1).

The preparation of the compound of formula B 2-(trimethylsilyl)-ethyl {(3S,4S)-3-amino-4-[(aminocarbonyloxy)methyl]-2-oxoazetidinyloxy}acetate was prepared as followed:

Preparation of 2-[(tert-butoxy)carbonylamino]-3,4-dihydroxy-N-(phenylmethoxy)butanamide

To solution of tetrahydrofuran (THF, 6 L) and phosphoric acid buffer (0.025 M, $\text{KH}_2\text{PO}_4/\text{Na}_2\text{HPO}_4$; ratio 1/1, 2 L) containing ethyl 3-[(tert-butoxy)carbonylamino]-2-hydroxy-3-[N-(phenylmethoxy) carbamoyl]propanoate (compound prepared from diethyl tartrate according to the procedures described in Org. Synth., Coll. Vol. 1998, 220, J. Org. Chem. 1983, 3556 and U.S. Pat. No. 4,794,108) (120 g, 31.38 mmol), sodium borohydride (59.35 g, 156.9 mmol) was added portion wise at 0° C. over 1 h. The resulting mixture was stirred at 0° C. for an additional 1 h and at room temperature for 2 h. The mixture was cooled at 0° C. before addition of 1M aqueous solution containing H_3PO_4 until the pH reach 8. NaCl (100 g) was added to the mixture and the organic layer was separated. Extraction with ethyl acetate (3x1.5 L) was performed and the combined organic phases were washed with brine (1 L), dried over MgSO_4 and evaporated under vacuo. The residue as purified by column chromatography using hexane/acetone as eluent to get 50 g of white solid.

-ESI-MS spectrum: m/z: 339 [M-1]⁺.

Preparation of 2-[(tert-butoxy)carbonylamino]-3-hydroxy-N-(phenylmethoxy)-4-[(1,1,2,2-tetramethyl-1-silapropoxy)butanamide

[(Tert-butoxy)carbonylamino]-3,4-dihydroxy-N-(phenylmethoxy)butanamide (47 g, 138 mmol), imidazole (37.5 g, 552 mmol) and tert-butyldimethylsilyl chloride (57.7 g, 386.4 mmol) were stirred at 0° C. for 1 h and the room temperature for an additional 1 h in a mixture of dichloromethane (1.7 L) and dimethylformamide (17 mL). Water (250 mL) was added and after decantation, water (2x250 mL) and brine (250 mL) was used for washing the organic phases, which was dried over Na_2SO_4 . After evaporation of the solvent, the residue was purified by column chromatography using a mixture of ethyl acetate/hexane (1/4) as eluent to get 28 g of the desired product.

¹H-NMR (CDCl_3) δ : 0.06 (s, 6H), 0.88 (s, 9H), 1.42 (s, 9H), 3.65-3.70 (m, 1H), 3.75-3.85 (m, 2H), 4.05-4.15 (m, 1H), 4.90 (s, 2H), 5.65-5.75 (m, 1H), 7.30-7.50 (m, 5H).

Preparation of N-{(3S,4S)-2-oxo-1-(phenylmethoxy)-4-[(1,1,2,2-tetramethyl-1-silapropoxy)methyl]azetidin-3-yl}(tert-butoxy)carboxamide

A solution of THF (1.5 L) containing 2-[(tert-butoxy)carbonylamino]-3-hydroxy-N-(phenylmethoxy)-4-[(1,1,2,2-tetramethyl-1-silapropoxy)butanamide (9.8 g, 21.56 mmol), triphenylphosphine (17.5 g, 66.8 mmol) and diethyl azodicarboxylate (11.26 g, 64.7 mmol) was stirred at room temperature for 2 h. The solvent was evaporated in vacuo and the residue was purified under column chromatography using a mixture of ethyl acetate and hexane (1/4) as eluent to obtained 7.25 g of the desired product.

¹H-NMR (CDCl_3) δ : 0.09 (s, 6H), 0.89 (s, 9H), 1.40 (s, 9H), 3.40-3.45 (m, 1H), 3.62 (d, 1H, J=7.2 Hz), 3.88 (d, 1H, J=7.2 Hz), 4.83 (m, 1H), 4.94 (s, 2H), 5.51 (d, 1H, J=6.5 Hz), 7.35-7.45 (m, 5H).

Preparation of N-[(3S,4S)-4-(hydroxymethyl)-2-oxo-1-(phenylmethoxy)azetidin-3-yl](tert-butoxy)carboxamide

N-{(3S,4S)-2-oxo-1-(phenylmethoxy)-4-[(1,1,2,2-tetramethyl-1-silapropoxy)methyl]azetidin-3-yl}(tert-butoxy)carboxamide (2.80 g, 6.41 mmol) was dissolved in THF (50 mL) and pyridine (5 mL). Pyridine-hydrofluoride complex (4.0 mL, 2.6 eq of pyridine, 24 eq of HF) was added at -20° C. After 10 min, the mixture was warmed up at room tem-

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perature and stirred for 3.5 h. Phosphoric acid buffer (0.025 M, $\text{KH}_2\text{PO}_4/\text{Na}_2\text{HPO}_4$; ratio 1/1, 400 mL) was added and the resulting solution was extracted with ethyl acetate (3×100 mL). The organic phase was dried over Na_2SO_4 and the solvent was evaporated in vacuo. 2.04 g of the desired product was obtained and used for the next step.

$^1\text{H-NMR}$ (DMSO-d_6) δ : 1.38 (s, 9H), 3.50-3.60 (m, 2H), 3.95-4.05 (m, 1H), 4.70-4.80 (m, 1H), 4.94 (s, 2H), 5.00-5.10 (m, 1H), 7.20-7.30 (m, 1H), 7.35-7.50 (m, 5H).

Preparation of {(2S,3S)-3-[(tert-butoxy)carbonylamino]-4-oxo-1-(phenylmethoxy)azetidin-2-yl}methyl aminooate (reaction done according to procedures described in Chem., Eur. J. 2005, 1949)

In anhydrous dichloromethane (100 mL), N-[(3S,4S)-4-(hydroxymethyl)-2-oxo-1-(phenylmethoxy)azetidin-3-yl] (tert-butoxy)carboxamide (1.0 g, 3.1 mmol) and trichloroacetyl chloride (1.11 mL, 9.3 mmol) were stirred at 5° C. for 30 min. Then aluminium oxide (9.6 g) was added and the solvent was removed in vacuo. After 2 h at room temperature, the residue was taken up with ethyl acetate (40 mL) and stirred for 30 min. The eluent was concentrated and the residue was purified by column chromatography using ethyl acetate/hexane (1/2) as eluent to afford 0.895 g of the desired compound.

$^1\text{H-NMR}$ (DMSO-d_6) δ : 1.39 (s, 9H), 3.90-4.20 (m, 3H), 4.80-4.90 (m, 1H), 4.91 (s, 2H), 6.50-6.80 (broad band for NH_2 , 2H), 7.30-7.50 (m, 5H), 7.55-7.60 (m, 1H).

Preparation of {(2S,3S)-3-[(tert-butoxy)carbonylamino]-1-hydroxy-4-oxoazetidin-2-yl}methyl aminooate

{(2S,3S)-3-[(tert-butoxy)carbonylamino]-4-oxo-1-(phenylmethoxy)azetidin-2-yl}methyl aminooate (0.086 g, 0.24 mmol) was dissolved in a mixture of ethyl acetate (4 mL) and methanol (4 mL) under hydrogen atmosphere at room temperature in presence of Pd/C (10%, 25 mg). The reaction was stirred for 2 h and the mixture was filtrated over celite bed. The filtrate was evaporated in vacuo to give 0.055 g of the desired compound.

$^1\text{H-NMR}$ (DMSO-d_6) δ : 1.39 (s, 9H), 3.95-4.10 (m, 3H), 4.75-4.80 (m, 1H), 6.45-6.70 (broad band for NH_2 , 2H), 7.57 (d, 1H, $J=6.0$ Hz).

Preparation of (trimethylsilyl)-ethyl 2-[(3S,4S)-3-amino-4-[(aminocarbonyloxy)methyl]-2-oxoazetidin-2-yl]acetate

The titled compound was prepared from {(2S,3S)-3-[(tert-butoxy)carbonylamino]-1-hydroxy-4-oxoazetidin-2-yl}methyl aminooate according to the procedures described in J. Med. Chem. 1985, 1447 and J. Antibiotics, 1985, 813.

$^1\text{H-NMR}$ (CDCl_3) δ : 0.02 (s, 9H), 0.99 (t, 2H, $J=8.6$ Hz), 1.37 (s, 9H), 3.90-4.30 (m, 5H), 4.53 (dd, 2H, $J=16.4$, 21.6 Hz), 4.84 (dd, 1H, $J=5.4$, 9.4 Hz), 6.70-6.40 (broad band for NH_2 , 2H), 7.48 (d, 1H, $J=9.6$ Hz).

Final assembly and deprotection steps were done similarly according to the method described for the example 1 and J. Med. Chem. 1985, 1447.

Example 13

((2S,3S)-3-[(2Z)-2-(5-amino(1,2,4-thiadiazol-3-yl))-3-[(1,5-dihydroxy-4-oxo(2-hydropyridyl)methoxy]-3-azaprop-2-enoylamino]-1-(hydroxysulfonyloxy)-4-oxoazetidin-2-yl)methyl aminooate (48)

The titled compound was prepared following scheme 1.

The preparation of the compound of formula A (2Z)-3-[[1,5-bis(diphenylmethoxy)-4-oxo(2-hydropyridyl)]meth-

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oxy]-2-{5-[(triphenylmethyl)amino](1,2,4-thiadiazol-3-yl)}-3-azaprop-2-enoic acid was achieved according to the procedures described in example 1 from 2-oxo-2-{5-[(triphenylmethyl)amino](1,2,4-thiadiazol-3-yl)}acetic acid (EP-A-0 333 154 and GB-A-2102423).

The preparation of the compound of formula B [(2S,3S)-3-amino-1-(hydroxysulfonyloxy)-4-oxoazetidin-2-yl]methyl aminooate was achieved according to the procedures described in J. Antibiotics 1985, 1536 from {(2S,3S)-3-[(tert-butoxy)carbonylamino]-1-hydroxy-4-oxoazetidin-2-yl}methyl aminooate (example 12).

Final assembly and deprotection steps were done similarly according to the method described for the example 1 and J. Med. Chem. 1985, 1447.

Compound 8 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding hydroxylamine derivative ($\text{NH}_2\text{—O—R}_4$) was prepared in analogy to the procedure described in J. Med. Chem. 2004, 6349.

Compound 9 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding azetidinone ring of formula B was prepared in analogy to the procedures described in J. Am. Chem. Soc. 1980, 7076 and J. Org. Chem. 1982, 5160.

Compound 13 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding azetidinone ring or formula B was prepared in analogy to the procedures described in J. Antibiotics 1986, 76.

Compound 14 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding azetidinone of formula B was prepared in analogy to the procedures described in DE-A-3229439.

Compound 15 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding heterocycle of the carboxylic acid derivative of formula A was prepared in analogy to the procedures described in Russ. J. Org. Chem. 1995, 240 and J. Antibiotics 1983 1020.

Compound 16 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding heterocycle of the carboxylic acid derivative of formula A was prepared in analogy to the procedures described in Z. Chem 1975, 233 and J. Antibiotics 1983, 1020.

Compound 17 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding heterocycle of the carboxylic acid derivative of formula A was prepared in analogy to the procedures described in U.S. Pat. No. 4,394,504 and J. Antibiotics 1983, 1020.

Compounds 18 and 19 were prepared in analogy to the procedures described in WO-A-02/22613. The corresponding heterocycles of the carboxylic acid derivatives of formula A was prepared in analogy to the procedures described in J. Am. Chem. Soc. 1959, 2452, WO-A-95/33724 and J. Antibiotics 1983, 1020 using respectively the methyl hydrazine and hydrazine as starting material.

Compound 20 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding heterocycle of the carboxylic acid derivative of formula A was prepared in analogy to the procedures described in U.S. Pat. No. 4,394,504 but using 2-amino-6-picoline as starting material.

Compound 24 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding azetidinone ring of formula B was prepared in analogy to the procedures described in example 13 and J. Antibiotics 1983, 1201.

Compound 25 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding azetidi-

none ring of formula B was prepared in analogy to the procedures described in example 13.

Compound 27 was prepared in analogy to the procedures described in WO-A-02/22613. The corresponding hydroxylamine derivative (NH₂—O—R₄) was prepared according to the synthetic scheme B-3-II.

Compound 30 was prepared in analogy to the procedures described in example 7 using the same compound of formula B and the compound of formula A described in example 13.

Compound 31 was prepared in analogy to the procedures described in example 6 using the same compound of formula B and the compound of formula A described in example 12.

Compound 32 was prepared in analogy to the procedures described in example 6 using the same compound of formula B and the compound of formula A described in example 13.

Compound 35 was prepared in analogy to the procedures described in examples 4 and 6 for preparing the compound of formula B and of example 13 for preparing compound of formula A.

Compound 36 was prepared in analogy to the procedures described in example 4 using the same compound of formula B and the compound of formula A described in example 12.

Compound 37 was prepared in analogy to the procedures described in example 4 using the same compound of formula B and the compound of formula A described in example 13.

Compound 38 was prepared in analogy to the procedures described in example 6 and after reduction of the N-hydroxy group to NH (J. Am. Chem. Soc. 1980, 7076), the final sulfonylation was performed in analogy to example 3 to obtain the compound of formula B. Procedures of example 12 were used for preparing the compound of formula A.

Compound 39 was prepared in analogy to the procedures described in example 6 and after reduction of the N-hydroxy group to NH (J. Am. Chem. Soc. 1980, 7076), the final sulfonylation was performed in analogy to example 3 to obtain the compound of formula B.

Compound 40 was prepared in analogy to the procedures described in example 6 and after reduction of the N-hydroxy group to NH (J. Am. Chem. Soc. 1980, 7076), the final sulfonylation was performed in analogy to example 3 to obtain the compound of formula B. Procedures of example 13 were used for preparing the compound of formula A.

Compound 41 was prepared in analogy to the procedures described in example 12 for preparing the compound of formula B.

Compound 43 was prepared in analogy to the procedures described in example 12 for preparing the compound of formula B. Procedures of example 13 were used for preparing the compound of formula A.

Compound 44 was prepared in analogy to the procedures described in example 12 and J. Antibiotics 1983, 1201 for preparing the compound of formula B.

Compound 45 was prepared in analogy to the procedures described in example 12 and J. Antibiotics 1983, 1201 for preparing the compound of formula B. Procedures of example 12 were used for preparing the compound of formula A.

Compound 46 was prepared in analogy to the procedures described in example 12 and J. Antibiotics 1983, 1201 for preparing the compound of formula B. Procedures of example 13 were used for preparing the compound of formula A.

Compound 47 was prepared in analogy to the procedures described in example 12 for preparing the compound of formula B. Procedures of example 12 were used for preparing the compound of formula A.

In the following table X the analytical data for all compounds of formula (I) made are presented:

TABLE X

Compound Number (Example Number)	¹ H-NMR (DMSO-d ₆) δ in ppm	Mass m/z:
5 (8)	1.41 (d, 3H, J = 6.0 Hz), 3.70 (m, 1H), 3.5-6 (broad band for OH and NH ₂), 4.20 (s, 3H), 4.43 (dd, 1H, J = 2.56-7.6 Hz), 5.32 (s, 2H), 6.85 (s, 1H), 6.98 (s, 1H), 8.52 (s, 1H), 9.46 (d, 1H, J = 7.6 Hz).	
6 (1)	1.41 (d, 3H, J = 6.0 Hz), 3.62 (s, 3H), 3.67 (m, 1H), 4.43 (dd, 1H, J = 2.5, 7.7 Hz), 5.06 (s, 2H), 6.26 (s, 1H), 6.75 (s, 1H), 7.23 (s br, 2H), 7.45 (s, 1H), 9.40 (d, 1H, J = 7.7 Hz).	
7 (9)	1.42 (d, 3H, J = 6.1 Hz), 3.71 (qd, 1H, J = 2.4 and 6.1 Hz), 4.47 (dd, 1H, J = 2.4 and 7.8 Hz), 5.32 (s, 2H), 6.87 (s, 1H), 7.09 (s, 1H), 7.2-7.5 (Broad Band for NH ₂), 8.50 (s, 1H), 9.53 (d, 1H, J = 7.8 Hz).	
9	3.33 (m, 1H), 3.67 (m, 1H), 4.98 (m, 1H), 5.28 (s, 2H), 6.85 (s, 1H), 6.99 (s, 1H), 7.16 (Broad Band for NH ₂), 8.09 (s, 1H), 9.46 (d, 1H, J = 7.8 Hz).	
10 (10)	1.42 (d, 3H, J = 6.1 Hz), 1.99 (s, 3H), 3.67 (m, 1H), 3.79 (m, 2H), 4.23 (m, 2H), 4.43 (m, 1H), 6.86 (s, 1H), 9.15 (m, 1H), 9.93 (S broad, 1H).	
11 (11)	1.42 (d, 3H, J = 6.2 Hz), 3.74 (m, 1H), 4.48 (m, 1H), 5.41 (s, 2H), 6.94 (s, 1H), 7.20 (s, 1H), 8.29 (s, 1H), 9.59 (d, 1H, J = 7.7 Hz).	
12 (2)	1.39 (d, 3H, J = 6.2 Hz), 4.02 (m, 1H), 4.48 (m, 1H), 5.30 (s, 2H), 6.84 (s, 1H), 7.05 (s, 1H), 7.27 (br, s, 2H), 8.16 (s, 1H), 9.69 (d, 1H, J = 7.7 Hz).	
13	1.61 (d, 3H, J = 6.2 Hz), 4.43 (m, 1H), 4.87 (m, 1H), 5.30 (s, 2H), 6.89 (s, 1H), 6.99 (s, 1H), 7.27 (br, s, 2H), 8.11 (s, 1H), 9.57 (d, 1H, J = 7.7 Hz).	
14		518 (M ₄ + 1)
15	1.41 (d, 3H, J = 6.2 Hz), 3.65 (m, 1H), 4.44 (m, 1H), 5.34 (s, 2H), 7.06 (s, 1H), 8.17 (s, 1H), 9.47 (d, 1H, J = 7.7 Hz).	
16	1.41 (d, 3H, J = 6.2 Hz), 3.62 (m, 1H), 4.44 (m, 1H), 5.27 (s, 2H), 6.76 (s, 1H), 7.83 (s, 1H), 8.10 (broad s, 2H), 9.64 (d, 1H, J = 7.7 Hz).	
17	1.42 (d, 3H, J = 6.2 Hz), 3.10 (m, 1H), 4.47 (m, 1H), 5.31 (s, 2H), 6.10 (s, 1H), 8.86 (s, 1H), 7.90 (s, 1H), 9.72 (d, 1H, J = 7.7 Hz).	
18	1.42 (d, 3H, J = 6.2 Hz), 3.51 (m, 3H), 3.65 (m, 1H), 4.42 (m, 1H), 5.17 (s, 2H), 5.47 (s, 1H), 6.10 (s, 1H), 6.85 (s, 1H), 7.20 (br, s, 2H), 7.95 (s, 1H), 9.28 (d, 1H, J = 7.7 Hz).	
19	1.38 (d, 3H, J = 6.2 Hz), 3.68 (m, 1H), 4.39 (m, 1H), 5.17 (s, 1H), 5.31 (s, 2H), 6.78 (s, 1H), 7.89 (s, 1H), 9.24 (d, 1H, J = 7.7 Hz).	
20	1.43 (d, 3H, J = 6.2 Hz), 3.73 (m, 1H), 4.48 (m, 1H), 5.38 (s, 2H), 6.72 (d, 1H, J = 5.7 Hz), 6.83 (d, 1H, J = 5.7 Hz), 6.98 (s, 1H), 7.62 (t, 1H, J = 5.7 Hz), 8.04 (s, 1H), 9.54 (d, 1H, J = 7.7 Hz).	
21 (3)	1.43 (d, 3H), 3.67 (m, 1H), 4.45 (m, 1H), 5.47 (s, 2H), 6.56 (s, 1H), 7.05 (s, 1H), 8.10-8.20 (m, 3H), 9.50 (d, 1H, J = 7.7 Hz).	
22 (4)	1.41 (d, 3H, J = 6.2 Hz), 4.01 (m, 1H), 4.37 (m, 1H), 4.52 (AB, 2H, J = 16.4 Hz), 5.18 (s, 2H), 6.73 (s, 1H), 6.81 (s, 1H), 7.28 (br, s, 2H), 7.84 (s, 1H), 9.44 (d, 1H, J = 7.7 Hz).	
23 (5)	1.35-1.45 (m, 5H), 3.91 (m, 1H), 4.42 (m, 1H), 4.53 (m, 1H), 5.29 (s, 2H), 6.84 (s, 1H), 6.99 (s, 1H), 7.37 (br, s, 2H), 8.16 (s, 1H), 9.51 (d, 1H, J = 7.7 Hz).	
24	4.40-4.80 (m, 3H), 5.20 (m, 1H), 5.27 (s, 2H), 6.81 (s, 1H), 7.06 (s, 1H), 7.30 (br, s, 2H), 8.17 (s, 1H), 9.56 (d, 1H, J = 7.7 Hz).	
25	4.15 (m, 2H), 4.42 (m, 1H), 5.20-5.30 (m, 3H), 6.40-6.60 (br, s, 2H), 6.82 (s, 1H), 6.91 (s, 1H), 7.20-7.40 (br, s, 2H), 7.99 (s, 1H), 9.42 (d, 1H, J = 7.7 Hz).	
26 (6)	1.22 (d, 3H), 1.42 (d, 3H), 4.63 (d, 1H, J = 7.7 Hz), 5.23 (s, 2H), 6.81 (s, 1H), 7.00 (s, 1H), 7.27 (br, s, 2H), 8.08 (s, 1H), 9.59 (d, 1H, J = 7.7 Hz).	

TABLE X-continued

Compound Number (Example Number)	¹ H-NMR (DMSO-d ₆) δ in ppm	Mass m/z:
27	1.38 (m, 3H), 3.72 (m, 1H), 4.44 (m, 1H), 5.12 (s, 2H), 6.81 (s, 1H), 7.00 (s, 1H), 7.27 (br, s, 2H), 8.08 (s, 1H), 9.59 (d, 1H, J = 7.7 Hz)	
28		488 (M + 1)
29 (7)	1.31 (d, 3H, J = 6.2 Hz), 3.96 (m, 1H), 4.44 (m, 1H), 5.27 (s, 2H), 6.80 (s, 1H), 7.40-7.60 (br, s, 2H), 7.82 (m, 1H), 8.11 (m, 1H), 8.22 (m, 1H), 8.75 (m, 1H), 9.39 (d, 1H, J = 7.7 Hz)	
30	1.38 (d, 3H, J = 6.2 Hz), 3.90 (m, 1H), 4.46 (m, 1H), 5.21 (s, 2H), 6.69 (s, 1H), 7.74 (s, 1H), 8.15 (br, s, 2H), 9.68 (d, 1H, J = 7.7 Hz)	
31	1.22 (s, 3H), 1.42 (s, 3H), 4.58 (d, 1H, J = 7.7 Hz), 5.17 (s, 2H), 6.87 (s, 1H), 7.40 (br, s, 2H), 7.88 (s, 1H), 9.56 (d, 1H, J = 7.7 Hz)	
32	1.24 (s, 3H), 1.43 (s, 3H), 4.61 (d, 1H, J = 7.7 Hz), 5.38 (s, 2H), 7.09 (s, 1H), 8.16 (br, s, 2H), 8.21 (s, 1H), 9.63 (d, 1H, J = 7.7 Hz)	
33	1.23 (s, 3H), 1.48 (s, 3H), 4.49 (AB, 2H, J = 16.0 Hz), 4.70 (d, 1H, J = 7.7 Hz), 5.28 (s, 2H), 6.85 (s, 2H), 7.04 (s, 1H), 8.16 (s, 1H), 9.55 (d, 1H, J = 7.7 Hz)	
34	1.23 (s, 3H), 1.47 (s, 3H), 4.48 (AB, 2H, J = 16.0 Hz), 4.64 (d, 1H, J = 7.7 Hz), 5.26 (s, 2H), 7.04 (s, 1H), 7.42 (br, s, 2H), 8.16 (s, 1H), 9.49 (d, 1H, J = 7.7 Hz)	
35	1.24 (s, 3H), 1.48 (s, 3H), 4.49 (AB, 2H, J = 16.0 Hz), 4.68 (d, 1H, J = 7.7 Hz), 5.30 (s, 2H), 6.88 (s, 2H), 7.98 (s, 1H), 8.17 (br, s, 2H), 9.54 (d, 1H, J = 7.7 Hz)	
36	1.41 (d, 3H, J = 6.2 Hz), 3.90 (m, 1H), 4.42 (m, 1H), 4.52 (AB, 2H, J = 16.4 Hz), 5.28 (s, 2H), 7.00 (s, 1H), 7.44 (br, s, 2H), 8.11 (s, 1H), 9.42 (d, 1H, J = 7.7 Hz)	
37	1.42 (d, 3H, J = 6.2 Hz), 3.89 (m, 1H), 4.42 (m, 1H), 4.52 (AB, 2H, J = 16.4 Hz), 5.33 (s, 2H), 6.91 (s, 1H), 8.05 (s, 1H), 8.17 (br, s, 2H), 9.43 (d, 1H, J = 7.7 Hz)	
38	1.24 (s, 3H), 1.46 (s, 3H), 4.58 (d, 1H, J = 7.7 Hz), 5.17 (s, 2H), 6.87 (s, 1H), 7.42 (br, s, 2H), 7.86 (s, 1H), 9.45 (d, 1H, J = 7.7 Hz)	
39	1.25 (d, 3H), 1.49 (d, 3H), 4.64 (d, 1H, J = 7.7 Hz), 5.25 (s, 2H), 6.83 (s, 1H), 7.04 (s, 1H), 7.31 (br, s, 2H), 8.12 (s, 1H), 9.47 (d, 1H, J = 7.7 Hz)	
40	1.25 (s, 3H), 1.48 (s, 3H), 4.61 (d, 1H, J = 7.7 Hz), 5.35 (s, 2H), 7.04 (s, 1H), 8.10-8.15 (m, 3H), 9.46 (d, 1H, J = 7.7 Hz)	
41	3.50-3.60 (m, 2H), 4.00 (m, 1H), 4.23 (m, 1H), 4.36 (m, 1H), 4.50 (m, 2H), 5.20-5.30 (m, 3H), 6.50-6.70 (br, s, 2H), 6.83 (s, 1H), 6.95 (s, 1H), 7.27 (s, 2H), 8.06 (s, 1H), 9.35 (d, 1H, J = 7.7 Hz)	
42 (12)	3.50-3.60 (m, 2H), 3.95 (m, 1H), 4.20-4.40 (m, 2H), 4.45 (m, 2H), 5.17 (m, 1H), 5.27 (s, 2H), 6.50-6.70 (br, s, 2H), 6.92 (s, 1H), 7.38 (s, 2H), 7.98 (s, 1H), 9.33 (d, 1H, J = 7.7 Hz)	
43	3.50-3.60 (m, 2H), 3.91 (m, 1H), 4.27 (m, 1H), 4.30 (m, 1H), 4.49 (m, 2H), 5.22 (m, 1H), 5.28 (s, 2H), 6.53 (br, s, 2H), 6.83 (s, 1H), 7.95 (s, 1H), 8.13 (s, 2H), 9.32 (d, 1H, J = 7.7 Hz)	
47	4.06 (m, 1H), 4.27 (m, 1H), 4.40 (m, 1H), 5.12 (m, 1H), 5.25 (s, 2H), 6.40-6.60 (br, s, 2H), 6.98 (s, 1H), 7.40 (s, 1H), 8.03 (s, 1H), 9.34 (d, 1H, J = 7.7 Hz)	
48 (13)	4.07 (m, 1H), 4.27 (m, 1H), 4.42 (m, 1H), 5.15 (m, 1H), 5.36 (s, 2H), 6.30-6.50 (br, s, 2H), 7.03 (s, 1H), 8.13 (s, 1H), 8.17 (s, 1H), 9.36 (d, 1H, J = 7.7 Hz)	

Example 14

(1S,5R)-2-[2-(3-carbamoyl-6-methylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (111)

The titled compound was prepared following scheme 11.

a) Preparation of 5-carbamoyl-1-carboxymethyl-2-methyl-pyridinium bromide

A solution of 6-methyl nicotinamide (400 mg, 2.94 mmol, 1.0 eq) and bromoacetic acid (408 mg, 2.94 mmol, 1.0 eq) in DMF (10 mL) was stirred at room temperature for 6 days. The reaction mixture was monitored by LCMS. Solvent was then evaporated and the crude product was purified by preparative HPLC to afford 136 mg of the expected compound.

¹H-NMR (DMSO-d₆) δ (ppm): 2.80 (s, 3H); 5.66 (s, 2H); 8.13 (s, 1H); 8.24 (d, J=8.3, 1H); 8.53 (s, 1H); 8.92 (dd, J=1.8 and 8.3, 1H); 9.47 (s, 1H).

b) (1S,5R)-2-[2-(3-carbamoyl-6-methylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (111)

5-carbamoyl-1-carboxymethyl-2-methyl-pyridinium bromide (91 mg, 0.47 mmol, 1.0 eq) was added at room temperature to a stirred solution of (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (J. Med. Chem. 1998, 3961) (90 mg, 0.47 mmol, 1.0 eq) in DMSO (4 mL), followed by 1-hydroxybenzotriazol (69 mg, 0.52 mmol, 1.1 eq), dicyclohexylcarbodiimide (106 mg, 0.52 mmol, 1.1 eq) and diisopropylethylamine (96 μL, 0.56 mmol, 1.2 eq). After stirring overnight at room temperature, the reaction mixture was filtered. The mother liquid was evaporated and the crude product was dissolved in CH₂Cl₂ (4 mL) and filtered. The resulting solid was purified by preparative HPLC to afford 46 mg of the expected compound.

¹H-NMR (DMSO-d₆) δ (ppm): 1.82 (m, 1H); 2.42 (m, 1H); 2.73 (d, J=5.3, 3H); 3.25 and 3.55 (2m, 1H); 4.05 (m, 1H); 4.40 and 4.62 (2t, J=4.7, 1H); 5.19 and 5.31 (2d, J=4.3, 1H); 5.55-6.05 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.12 (d, J=7.0, 1H); 8.22 (d, J=8.3, 1H); 8.53 (d, J=10.0, 1H); 8.89 (dd, J=1.8 and 8.3, 1H); 9.36 (dd, J=1.8 and 10.0, 1H).

Example 15

(1S,5R)-2-[(2-[3-(N-methylcarbamoyl)pyridinium]acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (101)]

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available N-methyl nicotinamide as starting materials.

+ESI-MS spectrum: m/z: 289 [M+H—SO₃]⁺.

Example 16

(1S,5R)-2-[2-(4-aminopyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (103)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diaz-

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abicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-aminopyridine as starting materials.

+ESI-MS spectrum: m/z: 326 [M]⁺.

Example 17

(1S,5R)-2-(2-(2-isoquinolinium)acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (104)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available isoquinoline as starting materials.

-ESI-MS spectrum: m/z: 360 [M-1]⁺.

Example 18

(1S,5R)-2-[2-(4-carbamoylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (105)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available isonicotinamide as starting materials.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.40 (m, 1H); 3.20 and 3.50 (2m, 1H); 4.05 and 4.12 (2m, 1H); 4.43 and 4.60 (2t, J=4.7, 1H); 5.21 and 5.29 (2d, J=4.3, 1H); 5.55-6.00 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.27 (m, 1H); 8.46 (m, 2H); 8.65 (m, 1H); 9.12 (m, 2H).

Example 19

(1S,5R)-7-oxo-2-(2-(2-5,6,7,8-tetrahydroisoquinolinium)acetyl)-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (106)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 5,6,7,8-tetrahydroisoquinoline as starting materials.

+ESI-MS spectrum: m/z: 365 [M]⁺.

Example 20

(1S,5R)-2-[2-(3-aminopyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (107)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-aminopyridine as starting materials.

+ESI-MS spectrum: m/z: 326 [M]

Example 21

(1S,5R)-2-(2-{3-[N-(carbamoylmethyl)carbamoyl]pyridinium}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (108)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diaz-

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abicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and 3-[N-(carbamoylmethyl)carbamoyl]pyridine as starting materials.

+ESI-MS spectrum: m/z: 411 [M]⁺.

5 3-[N-(carbamoylmethyl)carbamoyl]pyridine was prepared by reacting the commercially available nicotinyl chloride hydrochloride with glycineamide hydrochloride.

Example 22

10 (1S,5R)-2-{2-[3-(N-cyclopropylcarbamoyl)pyridinium]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (109)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available nicotinyl chloride hydrochloride and cyclopropylamine as starting materials.

1H-NMR (DMSO-d₆): 0.60 (m, 2H); 0.78 (m, 2H); 1.77 (m, 1H); 2.40 (m, 1H); 2.92 (m, 1H); 3.23 and 3.50 (2m, 1H); 4.01 and 4.10 (2m, 1H); 4.41 and 4.61 (2t, J=4.7, 1H); 5.20 and 5.29 (2d, J=4.3, 1H); 5.58-6.05 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.30 (m, 1H); 8.95 (m, 1H); 9.01-9.13 (m, 2H); 9.36 (m, 1H).

Example 23

(1S,5R)-2-{2-[4-(dimethylamino)pyridinium]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (112)

The titled compound was prepared following scheme 11 and in analogy to example 14 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(dimethylamino)pyridine as starting materials.

1H-NMR (DMSO-d₆): 1.22 (s, 3H); 1.26 (s, 3H); 1.74 (m, 1H); 2.38 (m, 1H); 3.20-3.50 (m, 1H); 3.98 and 4.08 (2dd, J=8.6 and 11.2, 1H); 4.38 and 4.48 (2t, J=4.7, 1H); 5.00-5.45 (m, 3H); 7.02 and 7.07 (2d, J=7.9, 2H); 8.11 and 8.16 (2d, J=7.9, 2H).

Example 24

2-(2-{3-[N-((3S)pyrrolidin-3-yl)carbamoyl]pyridinium}acetyl)(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (122)

Preparation of sodium (1S,5R)-2-(2-bromoacetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate

A solution of sulphur trioxide.DMF complex (4.92 g, 32.10 mmol, 1.5 eq) in DMF (10 mL) was added at 0° C. to a stirred solution of (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F, 5.25 g, 21.40 mmol, 1.0 eq) in DMF (110 mL). After 5 hours stirring at 0° C., the reaction mixture was concentrated. The remaining oil was dissolved in a minimum amount of H₂O and the pH was adjusted to 6 with saturated NaHCO₃ solution. The mixture was then concentrated under reduced pressure to afford 8.3 g of the expected sodium (1S,5R)-2-(2-bromoacetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate as a brown oil.

Preparation of tert-butyl (3S)-3-(3-pyridylcarbonylamino)pyrrolidinecarboxylate

Nicotinoyl chloride hydrochloride (286 mg, 1.61 mmol, 1.0 eq) was added at room temperature to a stirred solution of

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(S)-3-amino-1-N-BOC-pyrrolidine (300 mg, 1.61 mmol, 1.0 eq) in CH_2Cl_2 (9 mL), followed by triethylamine (337 μL , 2.42 mmol, 1.5 eq). After stirring overnight at room temperature, the reaction mixture was extracted and the organic layer was dried over Na_2SO_4 , filtered and evaporated to afford 457 mg of the expected compound.

Preparation of 2-[2-[3-(N-{(3S)-1-[(tert-butyl)oxycarbonyl]pyrrolidin-3-yl}carbamoyl)pyridinium]acetyl]-(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt

A solution of (1S,5R)-2-(2-bromoacetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (200 mg, 0.64 mmol, 1.0 eq) and tert-butyl (3S)-3-(3-pyridylcarbonylamino)pyrrolidinecarboxylate (149 mg, 0.51 mmol, 0.8 eq) in DMF (2 mL) was stirred at room temperature for 3 days. The reaction was monitored by LCMS. Then DMF was evaporated to afford 330 mg the expected crude product which was directly used in the next step.

Preparation of 2-(2-[3-[N-{(3S)pyrrolidin-3-yl}carbamoyl]pyridinium]acetyl)-(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (122)

A solution of 2-[2-[3-(N-{(3S)-1-[(tert-butyl)oxycarbonyl]pyrrolidin-3-yl}carbamoyl)pyridinium]acetyl]-(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt in DMF (4 mL) was cooled to 0°C . before the addition of trifluoroacetic acid (729 μL , 9.45 mmol, 15.0 eq). After stirring overnight at room temperature, the reaction mixture was concentrated and the crude was purified by preparative HPLC to afford 37 mg of the expected compound.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.80 (m, 1H); 2.05 (m, 1H); 2.23 (m, 1H); 2.40 (m, 1H); 3.33 (m, 5H); 4.01 and 4.10 (2m, 1H); 4.13 and 4.61 (2t, $J=4.7$, 1H); 4.56 (m, 1H); 5.18 and 5.30 (2d, $J=4.1$, 1H); 5.60-6.05 (AB part of an ABX system, the X part being in the ^{15}N spectrum, 2H); 8.35 (m, 1H); 9.07 (m, 3H); 9.38 (m, 1H); 9.44 (s, 1H).

Example 25

(1S,5R)-2-[2-(3-carbamoylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (102)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available nicotinamide as starting materials.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.78 (m, 1H); 2.40 (m, 1H); 3.23 and 3.50 (2m, 1H); 4.03 and 4.13 (2dd, $J=8.6$ and 11.0 , 1H); 4.42 and 4.63 (2t, $J=4.7$, 1H); 5.20 and 5.32 (2d, $J=4.3$, 1H); 5.55-6.05 (AB part of a ABX system, the X part being in the ^{15}N spectrum, 2H); 8.18 (d, $J=3.7$, 1H); 8.33 (q, $J=6.2$ and 7.9 , 1H); 8.57 and 8.62 (2s, 1H); 9.02-9.12 (m, 2H); 9.42 (d, $J=6.5$, 1H).

Example 26

(1S,5R)-2-[2-(3,4-dicarbamoylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (110)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bro-

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moacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3,4-pyridine dicarboxamide as starting materials.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.74 (m, 1H); 2.40 (m, 1H); 3.25 and 3.49 (2m, 1H); 4.03 and 4.11 (2m, 1H); 4.41 and 4.62 (2t, $J=4.7$, 1H); 5.22 and 5.29 (2d, $J=4.3$, 1H); 5.55-6.00 (AB part of a ABX system, the X part being in the ^{15}N spectrum, 2H); 8.04 (m, 1H); 8.14 (d, $J=10.8$, 1H); 8.24 (m, 2H); 8.45 (d, $J=9.1$, 1H); 9.10 (m, 1H); 9.24 (2s, 1H).

Example 27

(1S,5R)-{2-[(2-(4-(isopropyl)pyridinium)acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (113)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 4-isopropyl pyridine as starting materials.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.28 and 1.30 (2d, $J=2.3$, 6H); 1.77 (m, 1H); 2.42 (m, 1H); 3.10-3.55 (m, 2H); 4.00 and 4.10 (2m, 1H); 4.40 and 4.60 (2t, $J=4.7$, 1H); 5.19 and 5.29 (2d, $J=4.2$, 1H); 5.45-5.90 (AB part of a ABX system, the X part being in the ^{15}N spectrum, 2H); 8.11 (m, 2H); 8.81 (d, $J=6.7$, 1H); 8.87 (d, $J=7.1$, 1H).

Example 28

(1S,5R)-2-[2-[3-(methoxycarbonyl)-5-methyl-pyridinium]acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (114)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available methyl-5-methylnicotinate as starting materials.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.77 (m, 1H); 2.41 (m, 1H); 2.58 (d, $J=5.9$, 3H); 3.24 and 3.49 (2m, 1H); 3.98 (d, $J=2.5$, 3H); 4.02 and 4.12 (2m, 1H); 4.41 and 4.61 (2t, $J=4.7$, 1H); 5.20 and 5.29 (2d, $J=4.3$, 1H); 5.55-6.00 (AB part of a ABX system, the X part being in the ^{15}N spectrum, 2H); 8.95 (m, 1H); 9.06 and 9.15 (2s, 1H); 9.41 and 9.45 (2s, 1H).

Example 29

(1S,5R)-2-[2-[3-(methoxycarbonyl)-2-methylpyridinium]acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (115)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 2-methyl nicotinic acid methyl ester as starting materials.

$^1\text{H-NMR}$ ($\text{DMSO}-d_6$): 1.82 (m, 1H); 2.40 (m, 1H); 2.81 (d, $J=2.8$, 3H); 3.29 and 3.57 (2m, 1H); 3.96 (s, 3H); 4.06 (m, 1H); 4.40 and 4.62 (2t, $J=4.8$, 1H); 5.20 and 5.34 (2d, $J=4.3$, 1H); 5.65-6.05 (AB part of a ABX system, the X part being in the ^{15}N spectrum, 2H); 8.17 (m, 1H); 8.91 (m, 1H); 9.08 (m, 1H).

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Example 30

(1S,5R)-2-{2-[3-(methoxycarbonyl)pyridinium]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (116)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3-carboxypyridine methyl ester as starting materials.

¹H-NMR (DMSO-d₆): 1.80 (m, 1H); 2.39 (m, 1H); 3.22 and 3.52 (2m, 1H); 3.98 (d, J=2.3, 3H); 4.01 and 4.12 (2m, 1H); 4.41 and 4.61 (2t, J=4.7, 1H); 5.20 and 5.29 (2d, J=4.2, 1H); 5.65-6.05 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.35 (m, 1H); 9.07 (m, 1H); 9.13 and 9.21 (2m, 1H); 9.57 and 9.62 (2s, 1H); 12.75 (br, 1H).

Example 31

(1S,5R)-7-oxo-2-[2-(4-propanoylpyridinium)acetyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (117)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 4-propionyl pyridine as starting materials.

¹H-NMR (DMSO-d₆): 1.13 (dt, J=2.5 and 7.1, 3H); 1.80 (m, 1H); 2.41 (m, 1H); 3.15-3.55 (m, 3H); 4.02 and 4.11 (2dd, J=8.8 and 11.1, 1H); 4.41 and 4.61 (2t, J=4.7, 1H); 5.20 and 5.29 (2d, J=4.3, 1H); 5.60-6.05 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.53 and 8.56 (2d, J=7.1, 2H); 9.11 and 9.18 (2d, J=7.1, 2H).

Example 32

(1S,5R)-2-[(2-(4-(aminothioxomethyl)pyridinium)acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (118)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 4-pyridine carbothioamide as starting materials.

¹H-NMR (DMSO-d₆): 1.79 (m, 1H); 2.39 (m, 1H); 3.24 and 3.49 (2m, 1H); 4.00 and 4.10 (2m, 1H); 4.41 and 4.61 (2t, J=4.7, 1H); 5.21 and 5.28 (2d, J=4.3, 1H); 5.55-5.95 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.28 (dd, J=6.8 and 10.8, 2H); 8.96 and 9.02 (2d, J=6.8, 2H); 10.31 and 10.74 (2br, 2H); 12.7 (br, 1H).

Example 33

(1S,5R)-2-(2-{3-[(ethoxycarbonyl)methyl]pyridinium}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (119)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound

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F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available ethyl-3-pyridyl acetate as starting materials.

¹H-NMR (DMSO-d₆): 1.22 (dt, J=0.9 and 7.1, 3H); 1.75 (m, 1H); 2.41 (m, 1H); 3.22 and 3.49 (2m, 1H); 4.05 (m, 3H); 4.15 (dq, J=0.9 and 7.1, 2H); 4.40 and 4.61 (2t, J=4.6, 1H); 5.20 and 5.30 (2d, J=4.3, 1H); 5.50-5.95 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.19 (m, 1H); 8.61 (m, 1H); 8.87 and 8.92 (2m, 1H); 8.89 and 8.97 (2s, 1H).

Example 34

(1S,5R)-7-oxo-2-{2-[3-(trifluoromethyl)pyridinium]acetyl}-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (120)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3-(trifluoromethyl)pyridine as starting materials.

¹H-NMR (DMSO-d₆): 1.80 (m, 1H); 2.40 (m, 1H); 3.24 and 3.51 (2m, 1H); 4.04 and 4.12 (2m, 1H); 4.42 and 4.62 (2t, J=4.7, 1H); 5.21 and 5.29 (2d, J=4.1, 1H); 5.65-6.05 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 8.47 (m, 1H); 9.17 (nit, 1H); 9.24 and 9.30 (2d, J=6.2, 1H); 9.67 and 9.76 (2s, 1H); 12.7 (br, 1H).

Example 35

(1S,5R)-2-[2-(3,4-dimethylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (121)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3,4-lutidine as starting materials.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.40 (d, J=3.7, 3H); 2.47 (m, 1H); 2.55 (d, J=3.6, 3H); 3.21 and 3.49 (2m, 1H); 4.06 (m, 1H); 4.40 and 4.60 (2t, J=4.7, 1H); 5.20 and 5.28 (2d, J=4.3, 1H); 5.40-5.85 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 7.97 (m, 1H); 8.70 (m, 2H).

Example 36

(1S,5R)-7-oxo-2-{2-[3-benzylpyridinium]acetyl}-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (123)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3-benzylpyridine as starting materials.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.40 (m, 1H); 3.21 and 3.41 (2m, 1H); 4.01 and 4.10 (2m, 1H); 4.21 (d, J=7.3, 2H); 4.40 and 4.60 (2t, J=4.7, 1H); 5.20 and 5.28 (2d, J=4.2, 1H); 5.50-5.95 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 7.31 (m, 5H); 8.12 (m, 1H); 8.55 (t, J=7.6, 1H); 8.80 and 8.85 (2d, J=6.3, 1H); 8.90 and 9.00 (2s, 1H).

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Example 37

(1S,5R)-7-oxo-2-[2-(3-phenylpyridinium)acetyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (124)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3-phenyl pyridine as starting materials.

¹H-NMR (DMSO-d₆): 1.80 (m, 1H); 2.40 (m, 1H); 3.23 and 3.51 (2m, 1H); 4.10 (m, 1H); 4.42 and 4.62 (2t, J=4.7, 1H); 5.21 and 5.33 (2d, J=4.2, 1H); 5.60-6.00 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 7.62 (m, 3H); 7.88 (m, 2H); 8.28 (m, 1H); 8.93 (2d, J=6.3, 1H); 8.99 (m, 1H); 9.41 and 9.48 (2s, 1H).

Example 38

2-(2-{3-[N-((3R)pyrrolidin-3-yl)carbamoyl]pyridinium}acetyl)(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (125)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available nicotinoyl chloride hydrochloride and (R)-3-amino-1-N-BOC-pyrrolidine as starting materials.
+ESI-MS spectrum: m/z: 423 [M]⁺.

Example 39

(1S,5R)-2-[2-(4-amino-3-carbamoylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (126)

The titled compound was prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 4-amino-3-pyridinecarboxamide as starting materials.

¹H-NMR (DMSO-d₆): 1.75 (m, 1H); 2.40 (m, 1H); 3.22 and 3.45 (2m, 1H); 4.01 and 4.10 (2dd, J=8.6 and 11.3, 1H); 4.40 and 4.60 (2t, J=4.8, 1H); 4.95-5.50 (m, 3H); 7.03 (dd, J=7.4 and 9.4, 1H); 7.83 (br, 1H); 8.05 (m, 1H); 8.15 (br, 1H); 8.65 (m, 1H); 8.99 and 9.04 (2br, 2H).

Example 40

(1S,5R)-2-[2-(3-carbamoyl-5-methylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (127)

The titled compound has been prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 5-methyl nicotinamide as starting materials.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.39 (m, 1H); 2.54 (m, 3H); 3.23 and 3.49 (2m, 1H); 4.05 (m, 1H); 4.41 and 4.61 (2t, J=4.7, 1H); 5.20 and 5.29 (2d, J=4.3, 1H); 5.50-6.00 (AB part of a ABX system, the X part being in the ¹⁵N spectrum,

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2H); 8.52 (d, J=16.0, 1H); 8.89 (d, J=7.2, 1H); 8.95 and 9.04 (2s, 1H); 9.24 (m, 2H); 12.6 (br, 1H).

Example 41

(1S,5R)-2-[2-[3-(aminocarbonylamino)pyridinium]acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (128)

The titled compound has been prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and 3-pyridylcarbamide.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.40 (m, 1H); 3.23 and 3.49 (2m, 1H); 3.98 and 4.10 (2m, 1H); 4.40 and 4.59 (2t, J=4.7, 1H); 5.20 and 5.28 (2d, J=4.2, 1H); 5.40-6.00 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 6.51 (br, 2H); 8.00 (m, 1H); 8.28 (m, 1H); 8.49 (m, 1H); 9.20 (m, 1H); 9.67 (d, J=12.2, 1H).

3-Pyridylcarbamide was prepared according to the procedure described in Heterocycles 1983, 1899.

Example 42

(1S,5R)-2-[2-(5-amino-3-carbamoylpyridinium)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (129)

The titled compound has been prepared following scheme 11 and in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 5-amino-3-pyridinecarboxamide as starting materials.

¹H-NMR (DMSO-d₆): 1.78 (m, 1H); 2.41 (m, 1H); 3.22 and 3.46 (2m, 1H); 3.98 and 4.09 (2m, 1H); 4.40 and 4.59 (2t, J=4.6, 1H); 5.20 and 5.26 (2d, J=4.1, 1H); 5.40-5.85 (AB part of a ABX system, the X part being in the ¹⁵N spectrum, 2H); 6.88 (d, J=11.5, 2H); 7.95 (m, 1H); 8.07 (m, 2H); 8.41 (m, 2H).

Example 43

(1S,5R)-2-[N-(4-[(2-aminoethylamino)carbonylamino]phenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (324)

Preparation of (tert-butoxy)-N-{4-[(fluoren-9-ylmethoxy)carbonylamino]phenyl}carboxamide

Triethylamine (7.36 mL, 52.82 mmol, 1.1 eq) was added at 0° C. to a stirred solution of N-BOC-1,4-phenylene diamine (10.00 g, 48.02 mmol, 1.0 eq) in CH₃CN (240 mL), followed by 9-fluorenylmethyloxycarbonyl chloride (14.90 g, 57.62 mmol, 1.2 eq). The resulting mixture was allowed to come at room temperature. After 4 hours stirring at room temperature, the reaction mixture was filtered to afford 20.60 g of the crude expected product as a white powder which was used in the next step without any further purification.

¹H-NMR (DMSO-d₆): 1.46 (s, 9H); 4.29 (t, J=6.6, 1H); 4.44 (d, J=6.3, 2H); 7.30-7.45 (m, 8H); 7.75 (d, J=7.4, 2H); 7.91 (d, J=7.4, 2H); 9.22 (br, 1H); 9.59 (br, 1H).

Preparation of

N-(4-aminophenyl)(fluoren-9-ylmethoxy)carboxamide

TFA (55.30 mL, 717.76 mmol, 15.0 eq) was added at 0° C. to a stirred solution of (tert-butoxy)-N-{4-[(fluoren-9-yl-

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methoxy)carbonylamino]phenyl}carboxamide (20.60 g, 47.85 mmol, 1.0 eq) in CH_2Cl_2 (900 mL). The resulting solution was allowed to come at room temperature. After stirring overnight at room temperature, the reaction mixture was concentrated to dryness and the residue was triturated in water. Then the mixture was filtered to afford 15.80 g of the expected crude product as a white powder.

$^1\text{H-NMR}$ (DMSO-d_6): 4.30 (t, $J=6.4$, 1H); 4.49 (d, $J=6.4$, 2H); 7.06 (d, $J=7.7$, 2H); 7.40 (m, 6H); 7.74 (d, $J=7.4$, 2H); 7.91 (d, $J=7.4$, 2H); 8.95 (br, 2H); 9.73 (br, 1H).

Preparation of N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}(fluoren-9-ylmethoxy)carboxamide

N,N'-Disuccinimidylcarbonate (16.20 g, 63.26 mmol, 1.1 eq) was added at room temperature to a stirred solution of N-(4-aminophenyl)(fluoren-9-ylmethoxy)carboxamide (20.00 g, 60.53 mmol, 1.0 eq) in CH_3CN (1100 mL). After stirring overnight at room temperature, the reaction mixture was filtered to afford 28.50 g of the expected crude product as a white powder.

$^1\text{H-NMR}$ (DMSO-d_6): 2.83 (br, 4H); 4.31 (t, $J=6.4$, 1H); 4.48 (m, 2H); 7.20-7.50 (m, 8H); 7.5 (d, $J=7.4$, 2H); 7.91 (d, $J=7.4$, 2H); 9.72 (br, 1H); 10.67 (br, 1H).

Preparation of N-{4-[(2-[(tert-butoxy)carbonylamino]ethyl]amino)carbonylamino]phenyl}(fluoren-9-ylmethoxy)carboxamide

A solution of N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}(fluoren-9-ylmethoxy)carboxamide (16.10 g, 34.15 mmol, 1.0 eq) in $\text{H}_2\text{O}/\text{CH}_3\text{CN}$ (1/1, v/v, 360 mL) was reacted at room temperature with NaHCO_3 (2.86 g, 34.15 mmol, 1.0 eq) and N-BOC-ethylene diamine (5.47 g, 34.15 mmol, 1.0 eq). After stirring overnight at room temperature, the reaction mixture was filtered to afford 16.36 g of the expected crude product as a white solid.

$^1\text{H-NMR}$ (DMSO-d_6): 1.37 (s, 9H); 2.98 (m, 2H); 3.11 (m, 2H); 4.29 (t, $J=6.4$, 1H); 4.44 (d, $J=6.4$, 2H); 6.10 (m, 1H); 6.85 (m, 1H); 7.30-7.50 (m, 8H); 7.74 (d, $J=7.4$, 2H); 7.90 (d, $J=7.4$, 2H); 8.40 (s, 1H); 9.53 (br, 1H).

Preparation of N-(4-aminophenyl){2-[(tert-butoxy)carbonylamino]ethyl}amino)carboxamide

Piperidine (9.68 mL, 97.75 mmol, 5.0 eq) was added at room temperature to a stirred solution of N-(4-[(2-[(tert-butoxy)carbonylamino]ethyl)amino)carbonylamino]phenyl}(fluoren-9-ylmethoxy)carboxamide (10.10 g, 19.55 mmol, 1.0 eq) in DMF (140 mL). After 2 hours stirring at room temperature, water was added to the reaction mixture and precipitation occurred. The resulting mixture was filtered, and the liquid phase was concentrated to afford 6.75 g of the expected product as an orange oil:

$^1\text{H-NMR}$ (DMSO-d_6): 1.37 (s, 9H); 2.98 (m, 2H); 3.11 (m, 2H); 4.69 (s, 2H); 6.00 (t, $J=5.5$, 1H); 6.44 (d, $J=8.6$, 2H); 6.81 (t, $J=5.3$, 1H); 6.97 (d, $J=8.6$, 2H); 8.00 (s, 1H).

Preparation of {2-[(tert-butoxy)carbonylamino]ethyl}amino-N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}carboxamide

N,N'-Disuccinimidylcarbonate (5.49 g, 21.44 mmol, 1.1 eq) was added at room temperature to a stirred solution of N-(4-aminophenyl){2-[(tert-butoxy)carbonylamino]ethyl}amino)carboxamide (6.75 g, 19.49 mmol, 1.0 eq) in

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CH_3CN (350 mL). After stirring overnight at room temperature, the reaction mixture was filtered and to afford 9.70 g of the expected crude product as a light brown solid.

$^1\text{H-NMR}$ (DMSO-d_6): 1.37 (s, 9H); 2.82 (br, 4H); 2.99 (m, 2H); 3.11 (m, 2H); 6.12 (t, $J=5.2$, 1H); 6.85 (t, $J=5.5$, 1H); 7.27 (d, $J=8.9$, 2H); 7.36 (d, $J=8.9$, 2H); 7.95 (s, 1H); 8.53 (s, 1H).

Preparation of [(2-aminoethyl)amino]-N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}carboxamide

TFA (11.59 mL, 150.54 mmol, 5.0 eq) was added at room temperature to a stirred suspension of {2-[(tert-butoxy)carbonylamino]ethyl}amino-N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}carboxamide (13.8 g, 30.11 mmol, 1.0 eq) in CH_2Cl_2 (165 mL). After stirring overnight at room temperature, solvent was evaporated and the crude product was triturated with Et_2O to afford 14.2 g of the expected crude product as a beige solid and as the trifluoroacetic acid salt.

$^1\text{H-NMR}$ (DMSO-d_6): 2.82 (br, 4H); 2.88 (m, 2H); 3.30 (m, 2H); 6.51 (t, $J=5.6$, 1H); 7.30 (d, $J=8.9$, 2H); 7.40 (d, $J=8.9$, 2H); 7.77 (br, 3H); 8.85 (s, 1H); 10.61 (s, 1H).

Preparation of (1S,5R)-2-[N-(4-{[(2-aminoethyl)amino]carbonylamino}phenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (324)

(1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H, 2.0 g, 10.41 mmol, 1.0 eq) was dissolved in H_2O (12.5 mL). Then CH_3CN (100 mL) was added at room temperature to the solution, followed by NaHCO_3 (1.57 g, 18.73 mmol, 1.8 eq) and [(2-aminoethyl)amino]-N-{4-[(2,5-dioxoazolidinyloxy)carbonylamino]phenyl}carboxamide (6.89 g, 14.57 mmol, 1.4 eq). After stirring overnight at room temperature, the reaction mixture was filtered to afford 3.27 g of the expected (1S,5R)-2-[N-(4-{[(2-aminoethyl)amino]carbonylamino}phenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid as a white solid.

$^1\text{H-NMR}$ (DMSO-d_6): 1.65 (m, 1H); 2.30 (dd, $J=5.8$ and 13.5, 1H); 2.90 (m, 2H); 3.18 (m, 1H); 3.30 (m, 2H); 3.98 (m, 1H); 4.41 (t, $J=4.7$, 1H); 5.22 (d, $J=4.3$, 1H); 6.23 (t, $J=5.7$, 1H); 7.28 (d, $J=8.2$, 2H); 7.33 (d, $J=8.2$, 2H); 7.65 (br, 3H); 8.38 (s, 1H); 8.53 (s, 1H).

Example 44

Sodium (1S,5R)-2-[N-(3,4-dihydroxyphenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (306)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-amino-1,2-benzenediol as starting materials.

$^1\text{H-NMR}$ (DMSO-d_6): 1.64 (m, 1H); 2.29 (dd, $J=5.8$ and 13.6, 1H); 3.13 (m, 1H); 3.93 (dd, $J=8.3$ and 11.0, 1H); 4.37 (t, $J=4.8$, 1H); 5.20 (d, $J=4.3$, 1H); 6.50-6.70 (m, 2H); 6.97 (m, 1H); 7.84 and 8.15 (2s, 1H); 8.35 and 8.43 (2s, 1H); 8.77 and 8.82 (2s, 1H).

Example 45

Sodium (1S,5R)-2-[N-[3-(acetylamino)phenyl]carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (307)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diaz-

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abicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1-acetamido-3-aminobenzene as starting materials.

¹H-NMR (DMSO-*d*₆): 1.66 (m, 1H); 2.02 (s, 3H); 2.31 (dd, J=5.8 and 13.6, 1H); 3.17 (m, 1H); 3.98 (dd, J=8.3 and 11.0, 1H); 4.40 (t, J=4.8, 1H); 5.27 (d, J=4.3, 1H); 7.05-7.25 (m, 3H); 7.64 (m, 1H); 8.54 (s, 1H); 9.86 (s, 1H).

Example 46

Sodium (1S,5R)-7-oxo-2-[N-(3-sulfamoylphenyl)carbamoyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (308)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-aminobenzene sulfonamide as starting materials.

¹H-NMR (DMSO-*d*₆): 1.68 (m, 1H); 2.32 (dd, J=5.8 and 13.5, 1H); 3.21 (m, 1H); 4.02 (dd, J=8.5 and 11.2, 1H); 4.43 (t, J=4.7, 1H); 5.28 (d, J=4.2, 1H); 7.31 (br, 2H); 7.38-7.46 (m, 2H); 7.72 and 7.74 (2t, J=1.9, 1H); 8.05 (t, J=1.7, 1H); 8.89 (br, 1H).

Example 47

Sodium (1S,5R)-2-{N-[4-(dimethylamino)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (309)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(dimethylamino)aniline as starting materials.

+ESI-MS spectrum: m/z: 354 [M+H]

Example 48

Sodium (1S,5R)-2-[N-(4-{N-[2-(dimethylamino)ethyl]carbamoyl}phenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (310)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-aminobenzoic acid and 2-(dimethylamino)ethylamine as starting materials.

+ESI-MS spectrum: m/z: 425 [M+H].

Example 49

Sodium (1S,5R)-2-(N-{4-[N-(carbamoylmethyl)carbamoyl]phenyl}carbamoyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (312)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-aminobenzoic acid and glycine hydrochloride as starting materials.

¹H-NMR (DMSO-*d*₆): 1.68 (m, 1H); 2.32 (dd, J=5.8 and 13.5, 1H); 3.21 (m, 1H); 3.78 (d, J=5.7, 2H); 4.02 (dd, J=8.3 and 11.1, 1H); 4.42 (t, J=4.7, 1H); 5.28 (d, J=4.6, 1H); 7.02

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(br, 1H); 7.23 (br, 1H); 7.58 (d, J=9.1, 2H); 7.79 (d, J=9.1, 2H); 8.50 (t, J=5.8, 1H); 8.78 (s, 1H).

Example 50

Sodium (1S,5R)-2-[N-(3-(1,3-oxazol-5-yl)phenyl)carbamoyl]-7-oxo-2,5-diazabicyclo[3.2.0]heptane-6-sulfonate (318)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-(1,3-oxazol-5-yl)aniline as starting materials.

¹H-NMR (DMSO-*d*₆): 1.68 (m, 1H); 2.33 (dd, J=5.8 and 13.7, 1H); 3.21 (m, 1H); 4.02 (dd, J=8.1 and 11.0, 1H); 4.42 (t, J=4.7, 1H); 5.27 (d, J=4.3, 1H); 7.30-7.40 (m, 2H); 7.52 (m, 1H); 7.60 (s, 1H); 7.91 (s, 1H); 8.44 (s, 1H); 8.72 (br, 1H).

Example 51

Sodium (1S,5R)-7-oxo-2-[N-(2-oxo(3-hydrobenzimidazol-5-yl))carbamoyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (319)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 5(6)-aminobenzimidazolone as starting materials.

¹H-NMR (DMSO-*d*₆): 1.65 (m, 1H); 2.30 (dd, J=5.8 and 13.5, 1H); 3.17 (m, 1H); 3.96 (dd, J=8.3 and 11.0, 1H); 4.39 (t, J=4.7, 1H); 5.22 (d, J=4.3, 1H); 6.78 (m, 1H); 6.95 (m, 1H); 7.22 (m, 1H); 8.07 and 8.37 (2s, 1H); 10.34 and 10.39 (2s, 1H); 10.46 and 10.50 (2s, 1H).

Example 52

Sodium (1S,5R)-2-{N-[3-(ethoxycarbonyl)phenyl]carbamoyl}-7-oxo diazabicyclo[3.2.0]heptane-6-sulfonate (320)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-aminobenzoic acid ethyl ester as starting materials.

¹H-NMR (DMSO-*d*₆): 1.32 (t, J=7.1, 3H); 1.68 (m, 1H); 2.32 (dd, J=5.8 and 13.5, 1H); 3.20 (m, 1H); 4.02 (dd, J=8.3 and 11.0, 1H); 4.31 (q, J=7.1, 2H); 4.42 (t, J=4.7, 1H); 5.27 (d, J=4.5, 1H); 7.39 (t, J=8.0, 1H); 7.56 (2dd, J=1.2 and 1.6, 1H); 7.82 (m, 1H); 8.14 (t, J=1.9, 1H); 8.81 (br, 1H).

Example 53

Sodium (1S,5R)-2-{N-[3-(hydroxymethyl)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (321)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-(hydroxymethyl)aniline as starting materials.

¹H-NMR (DMSO-*d*₆): 1.67 (m, 1H); 2.31 (dd, J=5.8 and 13.5, 1H); 3.18 (m, 1H); 3.99 (dd, J=8.3 and 11.4, 1H); 4.41

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(t, J=4.7, 1H); 4.45 (d, J=5.8, 2H); 5.15 (t, J=5.6, 1H); 5.27 (d, J=4.3, 1H); 6.90 (m, 1H); 7.18 (t, J=7.8, 1H); 7.39 (m, 1H); 7.45 (m, 1H); 8.52 (br, 1H).

Example 54

(1S,5R)-2-[N-[4-({[2-(2-aminoethoxy)ethyl]amino}carbonylamino)phenyl]carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (323)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and 2,2-oxydiethylamine dihydrochloride (Eur. J. Org. Chem. 2002, 3004) as starting materials.

¹H-NMR (DMSO-d₆): 1.65 (m, 1H); 2.30 (dd, J=5.8 and 13.5, 1H); 2.99 (m, 2H); 3.16 (m, 1H); 3.29 (m, 2H); 3.49 (t, J=5.7, 2H); 3.60 (t, J=5.2, 2H); 3.95 (m, 1H); 4.39 (t, J=4.7, 1H); 5.20 (d, J=4.3, 1H); 6.13 (t, J=5.7, 1H); 7.24 (d, J=8.2, 2H); 7.31 (d, J=8.2, 2H); 7.73 (br, 3H); 8.35 (s, 1H); 8.37 (s, 1H).

Example 55

(1S,5R)-7-oxo-2-[N-{4-[(4-piperidylamino)carbonylamino]phenyl}carbamoyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (325)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and 1-BOC-4-amino-piperidine hydrochloride as starting materials.

¹H-NMR (DMSO-d₆): 1.52 (m, 2H); 1.68 (m, 1H); 1.98 (m, 2H); 2.31 (dd, J=6.0 and 13.6, 1H); 3.00 (m, 2H); 3.17 (m, 1H); 3.23 (m, 3H); 3.72 (m, 1H); 3.97 (dd, J=8.3 and 11.0, 1H); 4.40 (t, J=4.8, 1H); 5.21 (d, J=4.2, 1H); 6.28 (d, J=7.5, 1H); 7.23 (d, J=9.2, 2H); 7.32 (d, J=9.2, 2H); 8.12 (s, 1H); 8.28 (br, 1H); 8.38 (s, 1H).

Example 56

(1S,5R)-7-oxo-2-[N-[4-(piperazinylcarbonylamino)phenyl]carbamoyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (326)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and 1-BOC-piperazine as starting materials.

¹H-NMR (DMSO-d₆): 1.65 (m, 1H); 2.32 (dd, J=5.7 and 13.6, 1H); 3.13 (m, 4H); 3.19 (m, 1H); 3.62 (m, 4H); 3.97 (dd, J=8.3 and 11.2, 1H); 4.40 (t, J=4.7, 1H); 5.31 (d, J=4.3, 1H); 7.28 (d, J=9.2, 2H); 7.36 (d, J=9.2, 2H); 8.42 (s, 1H); 8.58 (s, 1H); 8.63 (br, 1H).

Example 57

Sodium (1S,5R)-2-[N-(4-aminophenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (327)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diaz-

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abicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available N-BOC-1,4-phenylene diamine as starting materials.

+ESI-MS spectrum: m/z: 326 [M+H]⁺.

Example 58

Sodium (1S,5R)-2-[N-(2-carbamoylphenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (328)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 2-aminobenzamide as starting materials.

¹H-NMR (DMSO-d₆): 1.67 (m, 1H); 2.36 (dd, J=6.0 and 13.8, 1H); 3.23 (m, 1H); 3.97 (m, 1H); 4.42 (t, J=4.7, 1H); 5.24 (d, J=4.2, 1H); 7.00 (m, 1H); 7.45 (m, 1H); 7.72 (br, 1H); 7.79 (dd, J=1.2 and 7.9, 1H); 7.91 (s, 1H); 8.28 (br, 1H); 8.37 (dd, J=1.2 and 8.4, 1H).

Example 59

Sodium (1S,5R)-7-oxo-2-[N-(4-{2-[(phenylmethoxy)carbonylamino]acetylaminophenyl}carbamoyl)-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (329)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and N-carbobenzoxylglycine as starting materials.

¹H-NMR (DMSO-d₆): 1.66 (m, 1H); 2.31 (m, 1H); 3.18 (m, 1H); 3.78 (d, J=6.1, 2H); 3.98 (dd, J=8.0 and 11.1, 1H); 4.40 (t, J=4.8, 1H); 5.03 (s, 2H); 5.23 (d, J=4.5, 1H); 7.22-7.58 (m, 10H); 8.49 (s, 1H); 9.83 (s, 1H).

Example 60

(1S,5R)-2-[N-(4-{{(2-morpholin-4-ylethyl)amino}carbonylamino}phenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (330)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and N-(2-aminoethyl)morpholine as starting materials.

¹H-NMR (DMSO-d₆): 1.67 (m, 1H); 2.31 (dd, J=6.1 and 13.7, 1H); 3.00-3.25 (m, 5H); 3.40-3.75 (m, 6H); 3.97 (m, 3H); 4.41 (t, J=4.7, 1H); 5.22 (d, J=4.2, 1H); 6.28 (br, 1H); 7.28 (d, J=9.2, 2H); 7.33 (d, J=9.2, 2H); 8.39 (s, 1H); 8.50 (s, 1H); 9.47 (br, 1H).

Example 61

(1S,5R)-2-[N-(4-{{[(2-morpholin-4-ylethyl)amino]carbonylamino}phenyl}carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (331)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and

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the commercially available 1,4-benzene diamine, N-(2-aminoethyl)morpholine and ethylenediamine as starting materials.

¹H-NMR (DMSO-*d*₆): 1.66 (m, 1H); 2.31 (dd, J=6.1 and 13.7, 1H); 3.00-3.22 (m, 11H); 3.49 (m, 2H); 3.61 (m, 2H); 3.96 (m, 3H); 4.39 (t, J=4.7, 1H); 5.22 (d, J=4.5, 1H); 6.11 (t, J=5.8, 1H); 6.32 (m, 2H); 7.24 (d, J=9.1, 2H); 7.33 (d, J=9.1, 2H); 8.38 (d, J=4.4, 1H); 9.53 (br, 1H).

Example 62

(1S,5R)-2-(N-{4-[N-(2-aminoethyl)carbamoyl]phenyl}carbamoyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (332)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-aminobenzamide and 2-bromoethanamine as starting materials.

¹H-NMR (DMSO-*d*₆): 1.69 (m, 1H); 2.33 (dd, J=5.8 and 13.7, 1H); 2.97 (m, 2H); 3.22 (m, 1H); 3.48 (m, 2H); 4.01 (dd, J=8.2 and 11.0, 1H); 4.43 (t, J=4.7, 1H); 5.27 (d, J=4.5, 1H); 7.61 (d, J=9.1, 2H); 7.75 (m, 5H); 8.43 (m, 1H); 8.81 (s, 1H).

Example 63

Sodium (1S,5R)-2-(N-{4-[(tert-butoxy)carbonylamino]phenyl}carbamoyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (333)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available N-BOC-1,4-benzene diamine as starting materials.

¹H-NMR (DMSO-*d*₆): 1.46 (s, 9H); 1.64 (m, 1H); 2.30 (dd, J=5.8 and 13.5, 1H); 3.16 (m, 1H); 3.97 (dd, J=8.5 and 11.3, 1H); 4.39 (t, J=4.8, 1H); 5.22 (d, J=4.3, 1H); 7.25-7.35 (m, 4H); 8.41 (s, 1H); 9.17 (s, 1H).

Example 64

Sodium (1S,5R)-2-{N-[(3,4-dihydroxyphenyl)methyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (303)

The titled compound was prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and commercially available 3,4-dihydroxybenzylamine.

-ESI-MS spectrum: m/z: 356 [M-H]⁺.

Example 65

(1S,5R)-2-{N-[4-(morpholin-4-ylmethyl)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (334)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(morpholinomethyl)aniline as starting materials.

¹H-NMR (DMSO-*d*₆): 1.67 (m, 1H); 2.30 (dd, J=5.8 and 13.6, 1H); 3.12 (m, 2H); 3.25 (m, 3H); 3.60 (t, J=11.8, 2H);

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3.97 (m, 3H); 4.26 (br, 2H); 4.42 (t, J=4.7, 1H); 5.25 (d, J=4.3, 1H); 7.36 (d, J=8.2, 2H); 7.60 (d, J=8.2, 2H); 8.74 (br, 1H); 9.61 (br, 1H).

Example 66

(1S,5R)-2-[N-(4-morpholin-4-ylphenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (335)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-morpholinoaniline as starting materials.

¹H-NMR (DMSO-*d*₆): 1.67 (m, 1H); 2.32 (dd, J=5.8 and 13.6, 1H); 3.19 (m, 1H); 3.32 (br, 4H); 3.85 (br, 4H); 3.98 (m, 1H); 4.42 (t, J=4.7, 1H); 5.25 (d, J=4.3, 1H); 7.24 (br, 2H); 7.50 (d, J=8.6, 2H); 8.60 (br, 1H).

Example 67

(1S,5R)-2-[N-(3-morpholin-4-ylphenyl)carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (336)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-morpholino-4-ylaniline as starting materials.

¹H-NMR (DMSO-*d*₆): 1.67 (m, 1H); 2.32 (dd, J=5.8 and 13.6, 1H); 3.19 (m, 1H); 3.26 (br, 4H); 3.82 (br, 4H); 3.99 (m, 1H); 4.42 (t, J=4.7, 1H); 5.25 (d, J=4.3, 1H); 6.84 (d, J=7.5, 1H); 7.15 (d, J=8.1, 1H); 7.23 (t, J=8.1, 1H); 7.48 (br, 1H); 8.60 (br, 1H).

Example 68

(1S,5R)-7-oxo-2-{N-[3-(piperazinylmethyl)phenyl]carbamoyl}-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (337)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available tert-butyl 4-(3-aminobenzyl)piperazine-1-carboxylate as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

¹H-NMR (DMSO-*d*₆): 1.67 (m, 1H); 2.32 (dd, J=5.8 and 13.6, 1H); 2.58 (br, 4H); 3.09 (br, 4H); 3.19 (m, 1H); 3.49 (br, 2H); 3.98 (m, 1H); 4.41 (t, J=4.7, 1H); 5.25 (d, J=4.3, 1H); 6.89 (d, J=7.5, 1H); 7.21 (t, J=8.1, 1H); 7.37 (t, J=8.1, 1H); 7.52 (br, 1H); 8.44 (br, 2H); 8.55 (br, 1H).

Example 69

2-[N-((3S)pyrrolidin-3-yl)carbamoyl](1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (338)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and commercially available (S)-3-amino-1-N-BOC-pyrroli-

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dine as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

+ESI-MS spectrum: m/z: 305 [M+H]⁺.

Example 70

2-[N-(4-[[((3S)pyrrolidin-3-yl)amino]carbonylamino]phenyl)carbamoyl](1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (339)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 1,4-benzene diamine and also commercially available (S)-3-amino-1-N-BOC-pyrrolidine as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

¹H-NMR (DMSO-d₆): 1.65 (m, 1H); 1.82 (m, 1H); 2.17 (m, 1H); 2.31 (dd, J=5.8 and 13.6, 1H); 3.06 (dd, J=5.1 and 11.8, 1H); 3.19 (m, 2H); 3.29 (m, 2H); 3.96 (dd, J=8.3 and 11.1, 1H); 4.22 (m, 1H); 4.40 (t, J=4.7, 1H); 5.22 (d, J=4.3, 1H); 6.40 (d, J=6.0, 1H); 7.26 (d, J=9.2, 2H); 7.33 (d, J=9.2, 2H); 8.37 (s, 1H); 8.41 (s, 1H); 8.70 (br, 2H).

Example 71

(1S,5R)-2-{N-[4-(2-morpholin-4-ylethoxy)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (340)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(2-morpholin-4-ylethoxy)aniline as starting materials.

¹H-NMR (DMSO-d₆): 1.66 (m, 1H); 2.31 (dd, J=5.8 and 13.6, 1H); 3.18 (m, 3H); 3.54 (m, 4H); 3.70 (m, 2H); 3.96 (m, 3H); 4.29 (m, 2H); 4.41 (t, J=4.7, 1H); 5.21 (d, J=4.3, 1H); 6.93 (d, J=9.0, 2H); 7.41 (d, J=9.0, 2H); 8.44 (s, 1H); 9.81 (br, 1H).

Example 72

(1S,5R)-2-{N-[3-(2-morpholin-4-ylethoxy)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (341)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 3-(2-morpholin-4-ylethoxy)aniline as starting materials.

¹H-NMR (DMSO-d₆): 1.66 (m, 1H); 2.31 (dd, J=5.8 and 13.6, 1H); 3.18 (m, 3H); 3.54 (m, 4H); 3.70 (m, 2H); 3.98 (m, 3H); 4.29 (m, 2H); 4.41 (t, J=4.7, 1H); 5.23 (d, J=4.3, 1H); 6.61 (dd, J=1.9 and 8.1, 1H); 7.04 (dd, J=1.1 and 8.1, 1H); 7.19 (t, J=8.1, 1H); 7.38 (s, 1H); 8.57 (s, 1H); 9.81 (br, 1H).

Example 73

(1S,5R)-7-oxo-2-[N-(4-piperidylphenyl)carbamoyl]-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (342)

The titled compound has been prepared following scheme and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diaz-

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abicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(1-piperidino)aniline as starting materials.

¹H-NMR (DMSO-d₆): 1.40-2.00 (m, 7H); 2.30 (dd, J=5.8 and 13.5, 1H); 3.22 (m, 1H); 3.45 (br, 4H); 3.98 (dd, J=8.3 and 11.0, 1H); 4.42 (t, J=4.7, 1H); 5.24 (d, J=4.3, 1H); 7.50-7.75 (m, 4H); 8.85 (br, 1H); 10.80 (s, 1H).

Example 74

(1S,5R)-2-[N-(6-morpholin-4-yl(3-pyridyl))carbamoyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (343)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 6-morpholinopyridine-3-amine as starting materials.

¹H-NMR (DMSO-d₆): 1.70 (m, 1H); 2.34 (dd, J=5.8 and 13.5, 1H); 3.22 (m, 1H); 3.54 (m, 4H); 3.76 (m, 4H); 3.98 (dd, J=8.3 and 11.0, 1H); 4.42 (t, J=4.7, 1H); 5.19 (d, J=4.4, 1H); 7.38 (d, J=9.7, 1H); 8.06 (dd, J=2.4 and 9.7, 1H); 8.29 (d, J=2.4, 1H); 8.88 (br, 1H).

Example 75

(1S,5R)-2-{N-[4-(4-methylpiperazinyl)phenyl]carbamoyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (344)

The titled compound has been prepared following scheme 12 and in analogy to example 43 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and the commercially available 4-(4-methylpiperazino)aniline as starting materials.

¹H-NMR (DMSO-d₆): 1.65 (m, 1H); 2.31 (dd, J=5.8 and 13.6, 1H); 2.69 (m, 2H); 2.99 (m, 2H); 3.05-3.25 (m, 8H); 3.96 (dd, J=8.3 and 11.0, 1H); 4.39 (t, J=4.7, 1H); 5.31 (d, J=4.3, 1H); 6.91 (d, J=9.0, 2H); 7.35 (d, J=9.0, 2H); 8.36 (s, 1H).

Example 76

(1S,5R)-2-{2-[1-(dimethylamino)-2-oxohydropyrimidin-4-ylthio]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (203)

The titled compound was prepared following scheme 12 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and 2-[1-(dimethylamino)-2-oxohydropyrimidin-4-ylthio]acetic acid. The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

+ESI-MS spectrum: m/z: 404 [M+H]⁺.

2-[1-(dimethylamino)-2-oxohydropyrimidin-4-ylthio]acetic acid was prepared from the 3-(dimethylamino)-6-sulfanyl-3-hydropyrimidin-2-one (U.S. Pat. No. 4,348,518) and bromoacetic acid according to the procedures described in Russian J. Org. Chem. 2000, 761.

Example 77

(1S,3R)-7-oxo-2-{2-[4-(2-pyridiniumacetyl)amino]phenylthio]acetyl}-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (204)

The titled compound was prepared following scheme 13 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]

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heptan-6-one (compound E) and the 2-[4-(2-pyridylacetylaminophenylthio)acetic acid as starting materials. The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

+ESI-MS spectrum: m/z: 397 [M+H—SO₃]⁺.

Example 78

(1S,5R)-2-(2-{1-[2-(dimethylamino)ethyl](1,2,3,4-tetraazol-5-ylthio)}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (205)

The titled compound was prepared following scheme 13 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and 2-{1-[2-(dimethylamino)ethyl]-1,2,3,4-tetraazol-5-ylthio}acetic acid as starting materials. The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

2-{1-[2-(dimethylamino)ethyl]-1,2,3,4-tetraazol-5-ylthio}acetic acid was prepared from commercially available 1-(2-dimethylaminoethyl)-5-mercapto-1,2,3,4-tetrazole and bromo acetic acid according to the procedures described in Russian J. Org. Chem. 2000, 761

+ESI-MS spectrum: m/z: 406 [M+H]⁺.

Example 79

Sodium (1S,5R)-2-[2-(1-methyl(1,3,4-thiadiazol-2-ylthio))acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (202)

The titled compound was prepared following scheme 13 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and the commercially available 2-(5-methyl-1,3,4-thiadiazol-2-ylthio)acetic acid as starting materials. The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

—ESI-MS spectrum: m/z: 363 [M—H]⁺.

Example 80

(1S,5R)-2-[2-(4-{N-[2-(dimethylamino)ethyl]carbamoyl}phenylthio)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (207)

[5-(2-Dimethylamino-ethylcarbamoyl)phenylthio]acetic acid

To a suspension of commercially available 4-mercapto benzoic acid (1.54 g, 10 mmol) in water was added NaOH (0.88 g, 22 mmol). The resulting solution was stirred at room temperature for 30 min and then bromo acetic acid ethyl ester (1.67 g, 10 mmol) was slowly added to the previous solution. After stirring at room temperature for 2 hours, an aqueous solution containing HCl (1M) was added. The obtained precipitate was filtrated, washed with water and dried to afford 1.5 g of 4-ethoxycarbonylmethylthio benzoic acid.

—ESI-MS spectrum: m/z: 239 [M—H]⁺.

Then the corresponding acid chloride of 4-ethoxycarbonylmethylthio benzoic acid was prepared in analogy to the procedure described in Synthesis, 1985, 517 and the condensation of the commercially available 2-dimethylaminoethyl amine and the hydrolysis of the ester group were performed in analogy to the procedure described in Bioorg. Med. Chem. Lett. 2003, 1517.

+ESI-MS spectrum: m/z: 283 [M+H]⁺.

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(1S,5R)-2-[2-(4-{N-[2-(dimethylamino)ethyl]carbamoyl}phenylthio)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (207)

[5-(2-Dimethylamino-ethylcarbamoyl)phenylthio]acetic acid (144 mg, 0.49 mmol, 1.0 eq) was added at room temperature to a stirred solution of (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H, 100 mg, 0.49 mmol, 1.0 eq) in DMSO (4 mL), followed by O-(benzotriazo-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate (HBTU) (225 mg, 0.59 mmol, 1.2 eq) and triethylamine (83 μ L, 0.59 mmol, 1.2 eq). After 4 hours stirring at room temperature, DMSO was evaporated, the crude was treated with acetonitrile. The resulting mixture was filtered to afford the crude product as a yellow solid which was purified by preparative HPLC: 74 mg (30%).

¹H-NMR (DMSO-d₆): 1.70 (m, 1H); 2.32 (m, 1H); 2.83 (s, 6H); 3.14 and 3.38 (2m, 1H); 3.26 (m, 2H); 3.57 (m, 2H); 3.96-4.23 (m, 3H); 4.34 and 4.51 (2t, J=4.7, 1H); 5.16 and 5.31 (2d, J=4.3, 1H); 7.43 (dd, J=8.5 and 12.6, 2H); 7.78 (dd, J=2.4 and 8.5, 2H); 8.62 (m, 1H); 9.18 (br, 1H).

Example 81

(1S,5R)-2-[2-(5-{N-[2-(dimethylamino)ethyl]carbamoyl}(2-pyridylthio))acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (208)

The titled compound has been prepared following scheme and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and [5-(2-dimethylaminoethylcarbamoyl)pyridin-2-ylthio]acetic acid as starting materials.

¹H-NMR (DMSO-d₆): 1.66 (m, 1H); 2.33 (m, 1H); 2.82 (s, 3H); 2.85 (s, 3H); 3.15 and 3.48 (2m, 1H); 3.26 (m, 2H); 3.61 (m, 2H); 3.98-4.55 (m, 4H); 5.16 and 5.35 (2t, J=4.3, 1H); 7.48 (d, J=8.4, 1H); 8.02 (m, 1H); 8.75 (m, 1H); 8.85 (dd, J=1.8 and 7.5, 1H); 9.16 (br, 1H).

[5-(2-Dimethylamino-ethylcarbamoyl)pyridin-2-ylthio]acetic acid was prepared in analogy to the procedure described in Bioorg. Med. Chem. Lett. 2003, 1517 using bromo acetic acid ethyl ester and commercially available 2-dimethylaminoethyl amine.

Example 82

(1S,5R)-2-(2-{4-[N-(2-aminoethyl)carbamoyl]phenylthio}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (209)

The titled compound has been prepared following scheme 13 and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and [4-(2-tert-butoxycarbonylaminoethylcarbamoyl)phenylthio]acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

¹H-NMR (DMSO-d₆): 1.72 (m, 1H); 2.33 (m, 1H); 2.97 (t, J=6.0, 2H); 3.14 and 3.38 (2m, 1H); 3.48 (q, J=6.0, 2H); 3.95-4.24 (m, 3H); 4.34 and 4.51 (2t, J=4.8, 1H); 5.17 and 5.32 (2d, J=4.3, 1H); 7.42 (dd, J=8.5 and 13.1, 2H); 7.78 (dd, J=2.5 and 8.5, 2H); 8.55 (m, 1H).

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[4-(2-Tert-butoxycarbonylaminoethylcarbamoyl)phenylthio]acetic acid was prepared in analogy to the procedure described in example 80 using 4-ethoxycarbonylmethylthiobenzoic acid and commercially available (2-aminoethyl) carbamic acid tert-butyl ester.

Example 83

(1S,5R)-2-(2-{4-[N-(2-aminoethyl)-N-methylcarbamoyl]phenylthio}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (210)

The titled compound has been prepared following scheme 13 and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and (4-[(2-tert-butoxycarbonylaminoethyl)methylcarbamoyl]phenylthio)acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 441 [M-H]⁺

{4-[(2-Tert-butoxycarbonylaminoethyl)methylcarbamoyl]phenylthio}acetic acid was prepared in analogy to the procedure described in example 80 using 4-ethoxycarbonylmethylsulfanylbenzoic acid and commercially available (2-methylamino-ethyl)carbamic acid tert-butyl ester.

Example 84

(1S,5R)-2-[2-(4-{N-[2-(methylamino)ethyl]carbamoyl}phenylthio)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (211)

The titled compound has been prepared following scheme 13 and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and {4-[2-(tert-butoxycarbonylmethylamino)ethylcarbamoyl]phenylthio}acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 441 [M-H]⁺

{4-[2-(Tert-butoxycarbonylmethylamino)ethylcarbamoyl]phenylthio}acetic acid was prepared in analogy to the procedure described in example 80 using 4-ethoxycarbonylmethylthiobenzoic acid and commercially available (2-aminoethyl)methylcarbamic acid tert-butyl ester.

Example 85

(1S,5R)-7-oxo-2-{2-[4-(piperazinylcarbonyl)phenylthio]acetyl}-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (212)

The titled compound has been prepared following scheme 13 and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and 4-(4-carboxymethylthiobenzoyl)piperazine-1-carboxylic acid tert-butyl ester as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 453 [M-H]⁺

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4-(4-Carboxymethylsulfanylbenzoyl)piperazine-1-carboxylic acid tert-butyl ester was prepared in analogy to the procedure described in example 80 using 4-ethoxycarbonylmethylthiobenzoic acid and commercially available piperazine-1-carboxylic acid tert-butyl ester.

Example 86

(1S,5R)-2-{2-[4-(2-aminoethoxy)phenylthio]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (213)

The titled compound has been prepared following scheme and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and [4-(2-tert-butoxycarbonylaminoethoxy)phenylthio]acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 400 [M-H]⁺

[4-(2-Tert-butoxycarbonylaminoethoxy)phenylthio]acetic acid was prepared in analogy to the procedure described in J. Med. Chem. 2000, 721 using first bromo acetic ethyl ester and then the commercially available (2-bromoethyl)carbamic acid tert-butyl ester.

Example 87

(1S,5R)-2-(2-{5-[N-(2-aminoethyl)carbamoyl](2-pyridylthio)}acetyl)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (214)

The titled compound has been prepared following scheme and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and [4-(2-tert-butoxycarbonylaminoethylcarbamoyl)phenylthio]acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 428 [M-H]⁺

[4-(2-Tert-butoxycarbonylamino-ethylcarbamoyl)phenylthio]acetic acid was prepared in analogy to the procedure described in Bioorg. Med. Chem. Lett. 2003, 1517 using bromo acetic acid ethyl ester and commercially available (2-aminoethyl)carbamic acid tert-butyl ester.

Example 88

(1S,5R)-2-[2-(5-{N-[2-(methylamino)ethyl]carbamoyl}(2-pyridylthio))acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (215)

The titled compound has been prepared following scheme and in analogy to example 80 using (1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (compound H) and {4-[2-(tert-butoxycarbonylmethylamino)ethylcarbamoyl]phenylthio}acetic acid as starting materials. The final deprotection step, using trifluoroacetic acid, has been performed in analogy to example 1.

-ESI-MS spectrum: m/z: 442 [M-H]⁺

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{4-[2-(Tert-butoxycarbonylmethylamino)ethylcarbamoyl]phenylthio}acetic acid was prepared in analogy to the procedure described in Bioorg. Med. Chem. Lett. 2003, 1517 using bromo acetic acid ethyl ester and commercially available (2-methylaminoethyl)carbamic acid tert-butyl ester.

Example 89

(1S,5R)-2-{2-[3-carbamoylpyridyl-4]carbonylamino]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (402)

The titled compound was prepared following scheme 11 in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) which was first sulfonated (J. Org. Chem. 1982, 5160) and the commercially available 3,4-pyridine dicarboxamide as starting materials.

+ESI-MS spectrum: m/z: 398 [M+H]⁺.

Example 90

Sodium 2-[[4(4S)-2-(2-hydroxyphenyl)(4,5-dihydro-1,3-thiazolin-4-yl)]carbonyl](1S,5R)-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (404)

The titled compound was prepared following scheme 13 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and the (4S) 4,5-dihydro-2-(2-hydroxyphenyl)-4-thiazolecarboxylic acid (JP59141554). The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

-ESI-MS spectrum: m/z: 397 [M+H]⁺.

Example 91

Sodium (1S,5R)-2-{2-[(5-fluoro-2-oxohydropyrimidin-4-yl)amino]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (405)

The titled compound was prepared following scheme 11 in analogy to example 22 using (5S,1R)-4-(2-bromoacetyl)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound F) and the commercially available 4-amino-5-fluoro pyridine-2-one.

-ESI-MS spectrum: m/z: 361 [M+H]⁺

Example 92

(1S,5R)-2-[2-amino-2-(4-carbamoylphenyl)acetyl]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonic acid (406)

The titled compound was prepared following scheme 13 in analogy to example 14 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and 2-amino-2-(4-carbamoylphenyl)acetic acid, obtained according to the procedure described in Eur. J. Med. Chem. 2003, 289 from 4-[[tert-butoxycarbonylamino](methoxycarbonyl)methyl]benzoic acid (WO-A-2000/076970). The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

+ESI-MS spectrum: m/z: 368 [M+H]⁺.

Example 93

(1S,5R)-2-{2-[4-(imidazolylcarbonyl)-1-methylpiperazinyl]acetyl}-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate, inner salt (407)

The titled compound was prepared following scheme 13 in analogy to example 24 using (5S,1R)-4-(2-bromoacetyl)-4,

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7-diazabicyclo[3.2.0]heptan-6-one (compound F) and 1-(1H-imidazol-1-ylcarbonyl)-4-methyl-piperazine (Ind. J. Chem., Section B, 1987, 748).

+ESI-MS spectrum: m/z: 426 [M]⁺.

Example 94

Sodium (1S,5R)-2-[[4-(4-carbamoylphenyl)amino]carbonylamino]-7-oxo-2,6-diazabicyclo[3.2.0]heptane-6-sulfonate (408)

The titled compound was prepared following the procedure described for example 18 of U.S. Pat. No. 6,566,355 using (5S,1R)-4,7-diazabicyclo[3.2.0]heptan-6-one (compound E) and 4-[(3-phenyl-1,2-oxaziridin-2-yl)carbonylamino]benzamide. The resulting compound was then sulfonated according to the procedure described in J. Org. Chem. 1982, 5160.

-ESI-MS spectrum: m/z: 369 [M+H]⁺

Biological Evaluation

Antimicrobial activity of the compounds and of their combinations was determined against a selection of organisms according to the standard procedures described by the National Committee for Clinical Laboratory Standards (National Committee for Clinical Laboratory Standards (2000). Methods for Dilution Antimicrobial Susceptibility Tests for Bacteria That Grow Aerobically-Fifth Edition: Approved Standard M7-A5. NCCLS, Wayne, Pa., USA).

The compounds to be tested were dissolved in 100% DMSO or sterile broth according to their aqueous solubility and were diluted to the final reaction concentration (0.06-32 µg/mL) in microbial growth media (IsoSensiTest Broth+16 µg/mL 2,2'-bipyridyl). In all cases the final concentration of DMSO incubated with the bacteria is less than or equal to 1%. For estimation of the minimal inhibitory concentrations (MIC), 2-fold dilutions of compounds were added to wells of a microtitre plate containing 10⁶ bacteria/mL. Plates were incubated overnight at an appropriate temperature (30° C. or 37° C.) and optical densities assessed by eye. The MIC value is defined as the lowest compound concentration completely inhibiting visible growth of the test organism. When combinations between compounds of formula I with compounds of formula II and are evaluated the compounds of formula I are tested in dilution series as described above while the compounds of formula II and III-XIII were present in all wells at a constant concentration of 4 µg/mL each.

The MIC values (in mg/L) of representative compounds and of representative combinations including these compounds are listed in tables 3, 4 and 5. Table 3 lists the MIC values obtained for two representative antibiotics of formula I when combined either with a compound of formula II or with a compound of formula III-XIII in comparison to the activity obtained with a combination involving the antibiotic of formula I with a compound of formula II and a compound of formula III-XIII together. Table 4 lists the activity of representative compounds of formula I alone or in combination with compounds of formula II and formula III-XIII. Table 5 lists the MIC values obtained for representative compounds of formula II with selected compounds of formula I and formula III-XIII.

If in table 4 in the upper three rows a cell is left empty, then this means that in the combination of that column no compound of the category of that row was used.

TABLE 3

Activity of representative monobactam antibiotics alone and in representative combinations according to the invention											
Strain	compound of for- mula I	1	Aztreonam								
	compound of for- mula II		206	323					206	323	
	Supplementary b- lactamase Inhibitor	clavulanate	501	clavulanate	clavulanate	501				sulbactam	
Minimal Inhibitory Concentration (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L											
Enterobac-ter cloacae	B1102	>32	32	16	16	8	2	16	2	0.125	2
	ATCC13047/ CRO-R	>32	>32	>32	>32	8	>32	>32	>32	8	8
	M4018	32	>32	16	8	2	>32	>32	32	1	4
	MRW	>32	32	32	16	4	>32	32	32	1	2
Enterobac-ter aerogenes	Zayakosky	>32	>32	>32	>32	16	>32	>32	>32	16	8
	B1	32	>32	>32	16	4	16	>32	32	8	8

TABLE 4

Activity of combinations of representative antibiotics									
Antibiotic	Mero- penem	Cefe- pime	Amox- icillin	Anip- icillin	Aztre- onam	Aztre- onam	Aztre- onam	1	1
β -lactamase inhibitor of formula II						102	102		
β -lactamase inhibitor of formula V or VI			clav- ulanate	sul- bactam	clav- ulanate	sul- bactam			501
Bacterial strain					MIC (mg/L) of antibiotic at fixed Inhibitor concentration of 4 mg/L				
<i>Acinetobacter baumannii</i> J2	2	32	>32	32	>32	16	>32	>32	<0.06
<i>Pseudomonas aeruginosa</i> MK1184	2	32	>32	>32	>32	32	16	>32	>32
<i>Pseudomonas aeruginosa</i> 1973E	1	4	>32	>32	8	8	4	16	8
<i>Enterobacter cloacae</i> P99	0.13	2	>32	>32	>32	2	2	>32	16
<i>Klebsiella pneumoniae</i> CF104	<0.06	4	32	>32	>32	0.5	1	>32	0.5
<i>Serratia marcescens</i> S6	16	0.13	>32	>32	32	2	2	>32	4
<i>Stenotrophomonas maltophilia</i> LAC736	32	4	>32	32	>32	4	8	1	0.25

Antibiotic	1	1	1	1	1	1	1	1	1
β -lactamase inhibitor of formula II									
β -lactamase inhibitor of formula V or VI									
Bacterial strain									
<i>Acinetobacter baumannii</i> J2	>32	4	2	8	4	4	4	1	16
<i>Pseudomonas aeruginosa</i> MK1184	>32	4	32	8	8	8	>32	4	16
<i>Pseudomonas aeruginosa</i> 1973E	8	2	4	4	4	4	4	4	8
<i>Enterobacter cloacae</i> P99	32	4	8	32	4	16	8	4	4

TABLE 4-continued

Activity of combinations of representative antibiotics											
Antibiotic	1	2	2	20	20	20	20	20	20	1	2
<i>Klebsiella pneumoniae</i> CF104	0.5	1	1	0.5	<0.06	2	0.25	1	2		
<i>Serratia marcescens</i> S6	4	0.25	0.5	0.25	0.25	0.5	4	0.5	0.25		
<i>Stenotrophomonas maltophilia</i> LAC736	0.5	0.125	2	0.125	0.125	0.125	0.125	0.125	0.125		
Antibiotic	1	2	2	20	20	20	20	20	20	20	20
β -lactamase inhibitor of formula II	324		206					102			206
β -lactamase inhibitor of formula V or VI	clav-ulanate		clav-ulanate		501		clav-ulanate		clav-ulanate		clav-ulanate
Bacterial strain					MIC (mg/L) of antibiotic at fixed Inhibitor concentration of 4 mg/L						
<i>Acinetobacter baumannii</i> J2	32	4	16	>32	<0.06	>32	>32	>32	>32	>32	>32
<i>Pseudomonas aeruginosa</i> MK1184	8	8	16	>32	>32	>32	>32	8	>32	>32	>32
<i>Pseudomonas aeruginosa</i> 1973E	8	8	8	16	16	16	16	32	16	16	16
<i>Enterobacter cloacae</i> P99	4	>32	16	>32	>32	>32	>32	16	16	16	16
<i>Klebsiella pneumoniae</i> CF104	1	>32	8	>32	16	16	32	2	4	4	4
<i>Serratia marcescens</i> S6	1	>32	>32	4	2	2	2	0.5	0.5	0.5	0.5
<i>Stenotrophomonas maltophilia</i> LAC736	0.125	4	0.5	4	0.25	0.25	0.25	0.25	0.5	0.5	0.5

Antibiotic of formula I	5	5	6	6	6	6	6	6	6	28	28
β -lactamase inhibitor of formula II		102		102	315	206	102	102	102		102
β -lactamase inhibitor of formula V or VI		clav-ulanate		clav-ulanate	clav-ulanate	clav-ulanate	clav-ulanate	clav-ulanate	clav-ulanate		clav-ulanate
Bacterial strain					MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L						
<i>Acinetobacter baumannii</i> J2	>32	32	>32	>32	>32	32	>32	16	>32	>32	>32

TABLE 4-continued

Activity of combinations of representative antibiotics												
	>32	>32	>32	>32	>32	>32	>32	>32	>32	>32	>32	>32
<i>Pseudomonas aeruginosa</i> MK1184	>32	>32	>32	>32	>32	>32	>32	>32	>32	16	>32	32
<i>Pseudomonas aeruginosa</i> 1973E	>32	>32	>32	>32	>32	>32	>32	>32	>32	16	>32	16
<i>Enterobacter cloacae</i> P99	32	8	16	8	4	16	>32	>32	>32	>32	>32	32
<i>Klebsiella pneumoniae</i> CF104	>32	8	>32	<0.06	0.5	0.5	>32	>32	>32	32	>32	2
<i>Serratia marcescens</i> S6	8	1	4	0.5	4	4	>32	>32	>32	>32	8	16
<i>Stenotrophomonas maltophilia</i> 1AC736	16	4	8	4	2	4	>32	>32	>32	2	8	8
Antibiotic of formula I	21	21	21	21	21	21	21	21	21	21	12	12
β -lactamase inhibitor of formula II		102		102		323		323		324		102
β -lactamase inhibitor of formula V or VI			sul-bactam	clav-ulanate		sul-bactam		clav-ulanate		sul-bactam		sul-bactam
Bacterial strain												
<i>Actinobacter baumannii</i> J2	>32	2		>32		8		8		2	>32	4
<i>Pseudomonas aeruginosa</i> MK1184	>32	32		16		32		32		32	>32	4
<i>Pseudomonas aeruginosa</i> 1973E	8	16		8		16		16		16	4	8
<i>Enterobacter cloacae</i> P99	>32	32		32		8		8		8	>32	8
<i>Klebsiella pneumoniae</i> CF104	>32	2		<0.06		1		2		4	>32	2
<i>Serratia marcescens</i> S6	8	16		0.5		2		2		4	32	1
<i>Stenotrophomonas maltophilia</i> 1AC736	8	8		2		4		2		4	1	4

MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L

Antibiotic of formula I	Activity of combinations of representative antibiotics									
	12	12	12	12	12	9	9	29	29	29
β-lactamase inhibitor of formula II	102	103	323	324			102			
clav- ulanate		sul- bactam	sul- bactam	sul- bactam			clav- ulanate	clav- ulanate	sul- bactam	sul- bactam
inhibitor of formula V or VI										
Bacterial strain										
<i>Actinobacter</i>	>32	1	2	2		>32	>32	>32	2	2
<i>baumannii</i> J2	16	16	16	16		>32	4	>32	4	8
<i>Pseudomonas aeruginosa</i>										
MK1184	4	16	4	16		8	8	4	8	8
<i>Pseudomonas aeruginosa</i>										
1973E										
<i>Enterobacter cloacae</i> P99	16	32	8	8		>32	32	>32	8	8
<i>Klebsiella pneumoniae</i> CF104	0.25	4	4	4		>32	0.5	>32	2	2
<i>Serratia marcescens</i> S6	2	16	4	16		16	1	32	8	8
<i>Stenotrophomonas maltophilia</i> 1AC736	0.125	2	0.5	4		0.5	0.25	1	1	1
Antibiotic of formula I	22	22	22	22	22	22	23	23	23	23
β-lactamase inhibitor of formula II										
clav- ulanate										
inhibitor of formula V or VI										
Bacterial strain										
<i>Actinobacter</i>	>32	4	4	8		2	16	8	8	8
<i>baumannii</i> J2	8	4	4	4		4	8	8	8	8
<i>Pseudomonas aeruginosa</i>										
MK1184	4	4	4	4		4	8	8	16	16
<i>Pseudomonas aeruginosa</i>										
1973E										
<i>Enterobacter cloacae</i> P99	>32	8	4	4		4	>32	8	4	4
<i>Klebsiella pneumoniae</i> CF104	>32	0.25	0.25	0.25		4	>32	1	1	1

TABLE 4-continued

Activity of combinations of representative antibiotics									
	0.13	0.25	0.13	0.13	0.25	0.25	0.25	0.25	0.25
<i>Serratia marcescens</i> S6									
<i>Stenotrophomonas maltophilia</i> LAC736	0.5	1	1	1	1				not tested
<i>Acinetobacter baumannii</i> J51						not tested			
<i>Enterobacter aerogenes</i> Zayakosky 5						not tested			
Antibiotic of formula I	23	26	26	26	26	26	26	26	26
β -lactamase inhibitor of formula II	324		102		324		324		102
β -lactamase of inhibitor of formula V or VI									
Bacterial strain									
<i>Acinetobacter baumannii</i> J2	2	4	4	2	4	2	2	4	4
<i>Pseudomonas aeruginosa</i> MK1184	8	2	2	2	2	2	2	2	2
<i>Pseudomonas aeruginosa</i> 1973E	16	16	2	4	2	2	2	2	2
<i>Enterobacter cloacae</i> P99	8	16	2	2	1	2	2	1	1
<i>Klebsiella pneumoniae</i> CF104	2	32	0.25	1	0.25	1	1	16	16
<i>Serratia marcescens</i> S6	0.5	<0.06	≤ 0.06	<0.06	<0.06	<0.06	<0.06	≤ 0.06	≤ 0.06
<i>Stenotrophomonas maltophilia</i> LAC736	not tested	<0.06	0.25	0.25	0.25	0.25	0.25	0.25	0.25
<i>Acinetobacter baumannii</i> J51	not tested	4	4	not tested	2	not tested	not tested	2	2
<i>Enterobacter aerogenes</i> Zayakosky 5	not tested	16	4	not tested	4	not tested	not tested	8	8

MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L

TABLE 4-continued

Activity of combinations of representative antibiotics						
Antibiotic of formula I	26	26	26	26	26	26
β -lactamase inhibitor of formula II	324					
β -lactamase of inhibitor of formula V or VI		clavulanic acid	501	sul- bactam	tazo- bactam	BRL427158
Bacterial strain			MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L			
<i>Acinetobacter</i>	4	>32	0.5	2	4	0.5
<i>baumannii</i> J2	4	4	2	4	4	2
<i>Pseudomonas</i>						
<i>aeruginosa</i>	4	2	2	2	2	2
MK1184						
<i>Pseudomonas</i>						
<i>aeruginosa</i>	2	8	2	8	16	2
1973E						
<i>Enterobacter</i>	16	0.5	1	1	1	0.125
<i>cloacae</i> P99						
<i>Klebsiella pneu-</i>						
<i>moniae</i> CF104	≤ 0.06	≤ 0.06	≤ 0.06	≤ 0.06	≤ 0.06	≤ 0.06
<i>Serratia</i>						
<i>marcescens</i> S6	0.25	≤ 0.06	0.125	≤ 0.06	≤ 0.06	0.125
<i>Stenotro-</i>						
<i>phomonas</i>						
<i>mallophilii</i>						
LAC736						
<i>Acinetobacter</i>	2	1	1	2	4	32
<i>baumannii</i> J51						
<i>Enterobacter</i>	4	8	16	16	16	0.5
<i>aerogenes</i>						
Zayakosky 5						
Antibiotic of formula I	41	41	41	48	48	48
β -lactamase inhibitor of formula II						
β -lactamase			324		102	324
inhibitor of formula V or VI		clavulanic acid	clavulanic acid	>32	clavulanic acid	clavulanic acid
Bacterial strain			MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L			
<i>Acinetobacter</i>	16	16	4	>32	>32	>32
<i>baumannii</i> J2	4	4	4	4	2	2
<i>Pseudomonas</i>						
<i>aeruginosa</i>	4	4	8	8	4	8
MK1184						
<i>Pseudomonas</i>						
<i>aeruginosa</i>						

TABLE 4-continued

Activity of combinations of representative antibiotics							
Antibiotic of formula I	25	25	25	47	47	47	47
1973E							
<i>Enterobacter cloacae</i> P99	>32	16	16	>32	32	16	
<i>Klebsiella pneumoniae</i> CF104	>32	0.5	0.5	>32	2	4	
<i>Serratia marcescens</i> S6	1	0.5	0.5	8	2	2	
<i>Stenotrophomonas maltophilia</i> LAC736	1	2	1	4	4	2	
Antibiotic of formula I	25	25	25	47	47	47	47
β -lactamase inhibitor of formula II							
β -lactamase inhibitor of formula V or VI							
Bacterial strain							
<i>Acinetobacter baumannii</i> J2	>32	16	>32	>32	>32	>32	
<i>Pseudomonas aeruginosa</i> MK1184	>32	4	8	4	2	4	
<i>Pseudomonas aeruginosa</i> 1973E	16	4	8	8	4	8	
<i>Enterobacter cloacae</i> P99	>32	8	4	>32	16	32	
<i>Klebsiella pneumoniae</i> CF104	>32	0.5	0.5	>32	4	2	
<i>Serratia marcescens</i> S6	1	1	1	2	2	2	
<i>Stenotrophomonas maltophilia</i> LAC736	4	1	1	1	1	1	

MIC (mg/L) of antibiotic at fixed inhibitor concentration of 4 mg/L

TABLE 4-continued

Antibiotic of formula I	Activity of combinations of representative antibiotics					
	32	32	32	33	33	33
β -lactamase inhibitor of formula II	102	324		102	324	
β -lactamase inhibitor of formula V or VI	clavulanic acid	clavulanic acid		clavulanic acid	clavulanic acid	
Bacterial strain	>32	4	8	>32	16	8
<i>Acinetobacter</i>						
<i>baumannii</i> J2						
<i>Pseudomonas</i>	4	4	4	32	16	32
<i>aeruginosa</i>						
MK1184	16	8	8	32	16	16
<i>Pseudomonas</i>						
<i>aeruginosa</i>						
1973E	32	8	4	32	4	16
<i>Enterobacter</i>						
<i>cloacae</i> P99						
<i>Klebsiella pneu-</i>	>32	1	2	2	0.5	0.5
<i>montiae</i> CF104						
<i>Serratia</i>	0.5	0.25	0.25	0.5	0.25	0.25
<i>marcescens</i> S6						
<i>Stenotro-</i>	0.5	0.5	0.5	2	2	2
<i>phomonas</i>						
<i>malophilia</i>						
1AC736						

TABLE 5

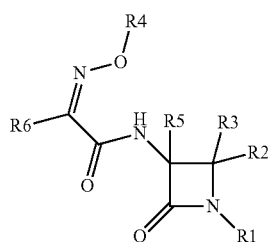
Activity of compounds of general formula II in representative combinations									
Bridged Mono-bactam of formula II	Supplementary β -lactamase Inhibitor of formula V or VI	MIC of antibiotic at fixed inhibitor concentration (4 mg/L)							
		Antibiotic of formula I	Acine-tobacter sp. J2	Pseudomonas aeruginosa MK1184	Pseudomonas aeruginosa 1973E	Enterobacter cloacae P99	Klebsiella pneumoniae CF104	Serratia marcescens S6	Stenotrophomonas maltophilia 1AC736
		Meropenem	2	2	1	0.125	<0.06	16	32
		Cefepime	32	32	4	2	4	0.125	4
		Aztreonam	>32	>32	8	>32	>32	32	>32
		1	>22	>32	16	>32	>32	1	0.5
101	clavulanate	1	>32	8	4	32	2	0.5	0.125
102	clavulanate	1	4	4	2	4	1	0.25	0.125
103	clavulanate	1	8	8	4	32	0.5	0.25	0.125
104	clavulanate	1	>32	32	4	32	0.5	2	0.125
105	clavulanate	1	8	16	8	32	0.5	0.125	0.125
106	clavulanate	1	>32	32	4	>32	0.25	0.25	0.125
107	clavulanate	1	8	8	4	32	1	0.125	0.125
108	clavulanate	1	>32	>32	4	4	1	0.25	0.125
109	clavulanate	1	4	16	4	32	0.5	0.125	0.125
110	clavulanate	1	2	32	4	>32	0.125	0.25	0.125
111	clavulanate	1	4	8	4	4	<0.06	0.25	0.125
112	clavulanate	1	16	>32	8	>32	<=0.06	0.5	0.125
113	clavulanate	1	>32	>32	4	32	0.125	0.25	0.25
114	clavulanate	1	>32	16	4	32	0.125	0.25	0.125
115	clavulanate	1	>32	16	4	32	0.125	0.25	0.125
116	clavulanate	1	16	8	16	32	0.5	0.25	0.125
117	clavulanate	1	>32	16	4	32	0.125	0.25	0.25
118	clavulanate	1	>32	32	4	32	0.125	0.25	0.125
119	clavulanate	1	>32	32	4	>32	0.125	0.25	0.125
120	clavulanate	1	32	16	8	32	0.25	0.25	0.125
121	clavulanate	1	>32	16	4	32	0.25	0.25	0.25
122	clavulanate	1	>32	16	4	16	0.125	0.25	0.125
123	clavulanate	1	>32	32	4	32	0.125	0.25	0.125
124	clavulanate	1	>32	32	9	32	0.125	0.25	0.125
125	clavulanate	1	4	16	8	32	<0.06	0.5	0.125
126	clavulanate	1	2	16	8	32	>32	0.125	0.125
201	clavulanate	1	4	32	8	32	>32	2	0.125
202	clavulanate	1	4	8	4	16	2	0.5	0.125
203	clavulanate	1	2	32	4	32	1	0.25	8
204	clavulanate	1	2	32	4	8	2	0.25	8
205	clavulanate	1	2	16	4	16	0.5	0.5	2
206	clavulanate	1	4	>32	4	8	0.25	4	0.125
301	clavulanate	1	8	>32	4	16	1	0.25	0.125
302	clavulanate	1	>32	>32	4	32	2	0.25	0.125
303	clavulanate	1	>32	>32	4	32	2	0.5	0.125
304	clavulanate	1	>32	>32	4	32	1	0.25	0.125
305	clavulanate	1	>32	>32	4	4	4	0.25	0.25
306	clavulanate	1	>32	>32	4	32	2	0.5	0.25
307	clavulanate	1	>32	>32	8	32	2	0.25	0.125
308	clavulanate	1	>32	>32	4	16	0.5	0.5	0.125
309	clavulanate	1	>32	>32	4	32	2	0.5	0.125
310	clavulanate	1	>32	16	4	4	2	0.5	0.125
311	clavulanate	1	4	>32	4	8	0.5	0.25	0.125
312	clavulanate	1	>32	>32	8	8	4	0.5	0.125
313	clavulanate	1	4	>32	4	16	0.5	0.25	0.125
314	clavulanate	1	4	>32	4	32	1	0.25	0.125
315	clavulanate	1	4	>32	4	8	1	0.25	0.125
316	clavulanate	1	16	32	4	8	0.5	0.25	0.125
317	clavulanate	1	16	>32	8	8	0.5	0.25	0.125
318	clavulanate	1	16	>32	4	>32	0.5	0.25	0.125
319	clavulanate	1	>32	>32	4	16	0.25	0.25	0.125
320	clavulanate	1	>32	>32	8	8	0.5	0.25	0.125
321	clavulanate	1	16	>32	4	>32	1	0.25	0.125
322	clavulanate	1	16	>32	8	>32	0.125	1	0.125
323	clavulanate	1	16	16	8	4	2	0.25	0.125
324	clavulanate	1	32	8	6	4	1	1	0.125
325	clavulanate	1	4	16	4	4	>32	0.25	0/25
326	clavulanate	1	4	32	4	8	>32	0.125	0.125
327	clavulanate	1	2	>32	2	32	2	0.25	0.125
328	clavulanate	1	8	32	4	4	1	1	0.125
331	clavulanate	1	>32	>32	4	4	0.5	2	0.125
332	clavulanate	1	>32	16	4	16	1	2	0.125
333	clavulanate	1	>32	32	4	32	1	3.25	0.125
401	clavulanate	1	4	>32	3	16	35	3.125	0.125
402	clavulanate	1	8	32	1	32	<0.06	0.25	<0.06
403	clavulanate	1	8	>32	2	32	4	0.5	0.125

TABLE 5-continued

Activity of compounds of general formula II in representative combinations									
Supplementary			MIC of antibiotic at fixed inhibitor concentration (4 mg/L)						
Bridged Mono-bactam of formula II	β -lactamase Inhibitor of formula V or VI	Antibiotic of formula I	Acinetobacter sp. J2	Pseudomonas aeruginosa MK1184	Pseudomonas aeruginosa 1973E	Enterobacter cloacae P99	Klebsiella pneumoniae CF104	Serratia marcescens S6	Stenotrophomonas maltophilia 1AC736
404	clavulanate	1	4	>32	4	16	0.5	0.25	0.25
406	clavulanate	1	>32	32	8	32	2	0.5	0.125
407	clavulanate	1	>32	>32	8	32	1	0.5	0.125
408	clavulanate	1	>32	>3	16	32	2	0.5	0.125

Further objects of the invention according to the following paragraphs 1-2 are also disclosed herein:

1. A pharmaceutical composition, comprising a combination of
a) an antibiotically active compound of the following formula I:



in which

R1 signifies SO_3H , OSO_3H , $\text{CRaRa}'\text{COOH}$, $\text{OCRaRa}'\text{COOH}$, 5-tetrazolyl, SO_2NHRb or CONHRc ,

wherein Ra and Ra' are independently selected from hydrogen; alkyl; allyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkylamino; dialkylamino; alkoxyalkyl and a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

wherein Rb is hydrogen; alkyl; alkoxy; carbonyl; alkylaminocarbonyl; benzylaminocarbonyl in which the benzyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or phenylaminocarbonyl in which the phenyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

wherein Rc is hydrogen; alkyl; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkoxy-carbonyl; SO_2phenyl ; $\text{SO}_2\text{NHalkyl}$; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

R2 and R3 independently signify hydrogen; alkyl; alkenyl; alkynyl; benzyl which may be substituted with 1 to 5 sub-

stituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; azido; halogen; dihalogenomethyl; trihalogenomethyl; alkoxy-carbonyl; carboxyl; sulfonyl or $\text{CH}_2\text{X1}$,

wherein X1 is azido; amino; halogen; hydroxyl; cyano; carboxyl; aminosulfonyl; alkoxy-carbonyl; alkanoylamino; phenylaminocarbonyl; alkylaminocarbonyl; aminocarbonyl; carbamoyloxy; alkylaminosulfonyl; phenylaminosulfonyl in which the phenyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

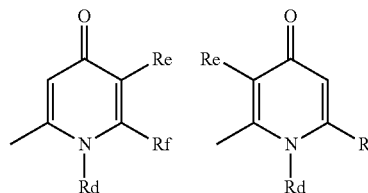
R4 signifies hydrogen; alkyl; C(Rx)(Ry)Z ,

wherein Rx and Ry are independently selected from hydrogen; alkyl; allyl; $(\text{C}_3\text{-C}_6)\text{cycloalkyl}$; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; $(\text{C}_2\text{-C}_7)\text{alkene}$ and $(\text{C}_2\text{-C}_7)\text{alkyne}$; or Rx and Ry taken together may form an alkylene bridge $\text{---}(\text{CH}_2)_n\text{---}$ with n being an integer number from 2 to 6; and

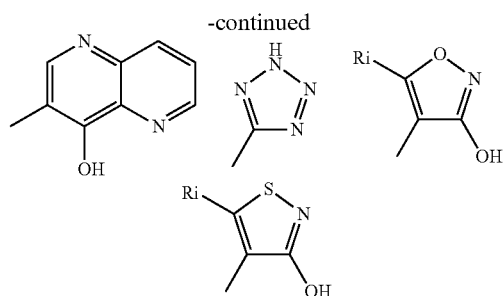
Z is COOH ; $\text{CH}_2\text{N(OH)COR}'$ wherein

R' is hydrogen, alkyl, alkylamino, alkoxy, benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen, phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen, or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

or Z is one of the following six groups



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in which groups

Rd, Re and Rf are independently selected from hydrogen; alkyl; amino; monoalkylamino; carboxylaminoalkyl; alkoxycarbonyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; diphenylmethyl; trityl; and ORg wherein

Rg is hydrogen; alkyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino and halogen; or phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino and halogen;

or, when Re and Rf are vicinal substituents, Re and Rf taken together may also be $-\text{O}-\text{CH}=\text{CH}-\text{CH}_2-$, $-\text{O}-\text{CH}_2-\text{CH}_2-\text{O}-$, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-$, $-\text{CH}=\text{CH}-\text{CH}=\text{CH}-$ or $-\text{CH}=\text{C}(\text{OH})-\text{C}(\text{OH})=\text{CH}-$;

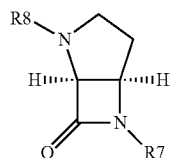
Ri is hydrogen; alkyl; alkylamino; alkoxy; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl and hydroxyl; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, dialkylamino and halogen;

R5 signifies hydrogen, alkyl, halogenomethyl, dihalogenomethyl, trihalogenomethyl, alkoxy, formylamino or alkylcarbonylamino;

R6 signifies phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, dialkylamino and halogen; or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, carbonylamino and halogen; or a pharmaceutically acceptable salt thereof;

with a β -lactamase inhibitor of one of the following groups b1) to b11):

b1) a bridged monobactam derivative of the following formula II:



in which:

R7 signifies SO_3H , OSO_3H or $\text{OCRjRj}'\text{COOH}$, wherein Rj and Rj' are independently selected from hydrogen; alkyl; phenyl which may be substituted with 1 to 5

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substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, dialkylamino and halogen; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, dialkylamino and halogen; alkylamino and alkoxyalkyl;

R8 is alkoxycarbonylamino, the acyl residue of an α or β -amino acid, or a residue of the formula $\text{Q}-(\text{X})_r-\text{Y}-$,

wherein Q is a 3-6 membered ring which optionally contains nitrogen, sulphur and/or oxygen and which is optionally fused to a phenyl ring or to a 5-6 membered heterocyclic ring and which is optionally substituted with 1 to 4 substituents selected from alkyl, allyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, dialkylamino, carboxamide which may be substituted, carboxylic acid, carbonylalkoxy, aminocarbonyl, alkylaminocarbonyl, halogen, halogenomethyl, dihalogenomethyl, trihalogenomethyl, sulfamide, substituted sulfamide with substituents selected from alkyl, allyl, phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino and halogen and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxyalkoxy, amino, alkylamino, halogen and benzyl, urea which may be substituted with alkyl, aminoalkyl, alkoxyalkyl or aminoalkoxyalkyl, and carbamate which may be substituted with alkyl, aminoalkyl, alkoxyalkyl or aminoalkoxyalkyl,

X signifies a linear spacer of from 1 to 6 atoms length and containing carbon, nitrogen, oxygen and/or sulphur atoms, of which up to 2 atoms can be nitrogen atoms and 1 atom can be oxygen or sulphur,

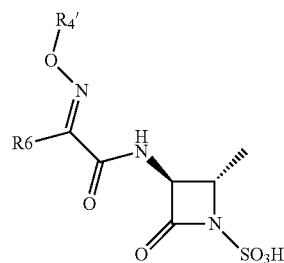
r is an integer of from 0 to 1; and

Y is selected from $-\text{CO}-$, $-\text{CS}-$, $-\text{NHCO}-$, $-\text{NHCONH}-$ and $-\text{SO}_2-$;

or a pharmaceutically acceptable salt thereof,

or

b2) a monobactam derivative of the general formula III:

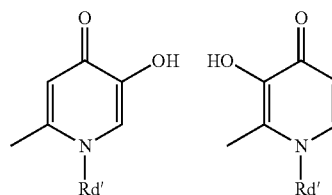


III

in which

R4' signifies hydrogen, alkyl or $\text{CH}(\text{Rx}')\text{Z}'$, wherein

Rx' is selected from hydrogen; (C_1-C_6) alkyl; allyl; phenyl and (C_3-C_6) cycloalkyl; and Z' signifies COOH or a group of one of the following two formulae:



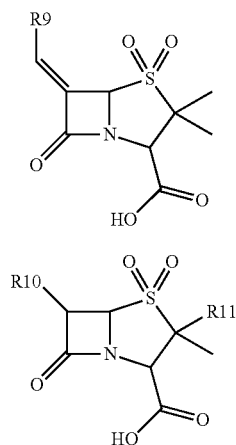
in which Rd' is hydrogen or hydroxy; and

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R6 is as defined for formula I; or a pharmaceutically acceptable salt thereof;

or

b3) a penam sulfone derivative of the general formulae IV or V;



in which

R9 signifies COOH or a 5-6 membered monocyclic or polycyclic heteroaromatic group;

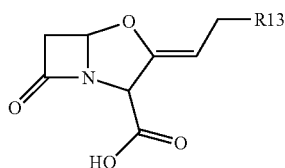
R10 signifies hydrogen or halogen;

R11 signifies CH_2R_{12} ; $\text{CH}=\text{CHR}_{12}$ wherein R12 is hydrogen, halogen, cyano, carboxylic acid, acyl such as acetyl, carboxamide which may be substituted, alkoxy-carbonyl or a 5-6 membered heteroaromatic ring which is optionally substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or which is optionally fused with a 5-6 membered heteroaromatic ring; $\text{CH}=\text{NR}_{12}'$ wherein R12' is amino, alkylamino, dialkylamino, aminocarbonyl, acylamino such as acetylamino, hydroxy, alkoxy,

or a pharmaceutically acceptable salt thereof;

or

b4) an oxapenam derivative of the general formula VI:



in which

R13 signifies OR_{14} ; $\text{S(O)}_n\text{R}_{14}$ or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; whereby $n=0, 1$ or 2 , and R14 is hydrogen, alkyl, $(\text{C}_2\text{-C}_7)\text{alkene}$, $(\text{C}_2\text{-C}_7)\text{alkyne}$ or a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents

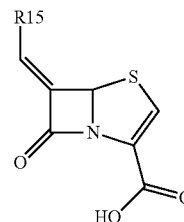
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selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen,

or a pharmaceutically acceptable salt thereof;

or

b5) a penem derivative of the general formula VII:



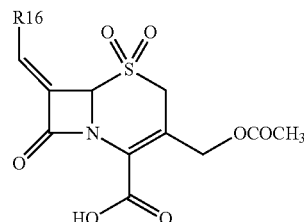
in which

R15 signifies a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or which is optionally fused with a 5-6 membered heteroaromatic ring and/or which is optionally bound to the exo-methylene group over a —CH=CH— spacer being preferably in the (E)-configuration,

or a pharmaceutically acceptable salt thereof;

or

b6) a cephem sulfone derivative of the general formula VIII:



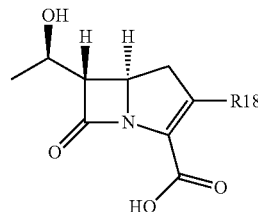
in which

R16 signifies COOR_{17} , whereby R17 signifies hydrogen or alkyl; or a 5-6 membered heteroaromatic ring which is optionally fused with a 5-6 membered heteroaromatic ring being optionally substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino, halogen; and/or being optionally bound to the exo-methylene group over a —CH=CH— spacer being preferably in the (E)-configuration,

or a pharmaceutically acceptable salt thereof;

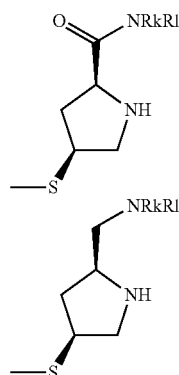
or

b7) a carbapenem derivative of the general formula IX:



in which R18 signifies —S-alkyl , $\text{—S—(CH}_2\text{)}_2\text{—NH—CH=NH}$ or a group of the following two formulae

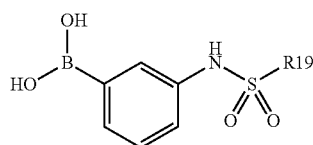
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wherein Rk and RI are individually selected from hydrogen, alkyl, 2-, 3-, 4-carboxyphenyl and sulfamoyl, or a pharmaceutically acceptable salt thereof;

or

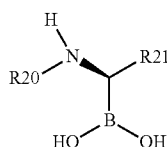
b8) a boronate derivative of the general formula X:



wherein R19 signifies a 5-6 membered heteroaromatic ring which may be substituted with amino, alkylamino, dialkylamino or alkylsulfoxide, or a pharmaceutically acceptable salt thereof;

or

b9) a boronate derivative of the general formula XI:



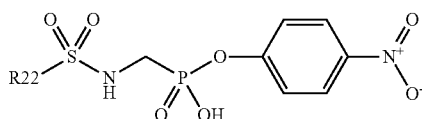
wherein

R20 and 21 are independently selected from a 5-6 membered heteroaromatic ring; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen,

or a pharmaceutically acceptable salt thereof;

or

b10) a phosphonate derivative of the general formula XII:



wherein

R22 is selected from a 5-6 membered heteroaromatic ring which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alky-

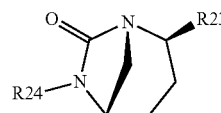
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lamino, dialkylamino and halogen and which is optionally fused with a 5-6 membered heteroaromatic ring; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; and benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen,

or a pharmaceutically acceptable salt thereof;

or

b11) a diazabicyclooctane derivative of the general formula XIII:



XIII

X in which

R23 signifies hydrogen, carboxylic acid, alkoxycarbonyl or carboxamide which may be substituted, and

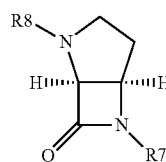
R24 signifies SO₃H, OSO₃H or OCR_jR_j'COOH, wherein R_j and R_j' are as defined for formula II,

or a pharmaceutically acceptable salt thereof.

2. Any embodiment as disclosed in the following claims 2 to 45, with their dependency or reference to claim 1 being replaced by a reference to preceding paragraph 1.

The invention claimed is:

1. A β -lactamase inhibitor of formula II:



II

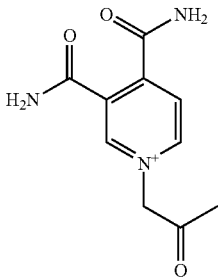
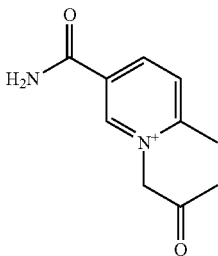
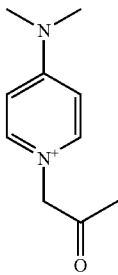
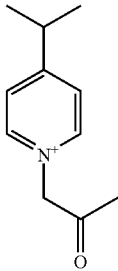
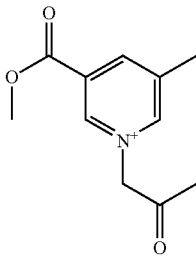
wherein

R7 and R8 are selected from the following table:

Compound number	R7	R8
106	SO ₃ ⁻	

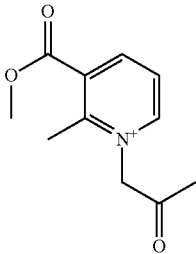
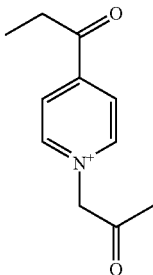
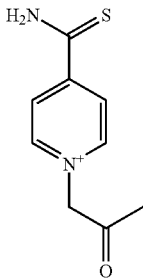
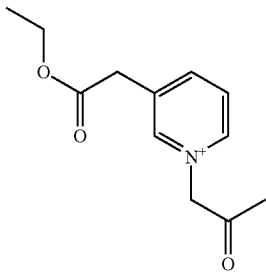
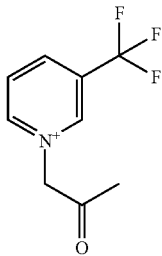
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-continued

Compound number	R7	R8
110	SO ₃ ⁻	
111	SO ₃ ⁻	
112	SO ₃ ⁻	
113	SO ₃ ⁻	
114	SO ₃ ⁻	

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-continued

Compound number	R7	R8
115	SO ₃ ⁻	
117	SO ₃ ⁻	
118	SO ₃ ⁻	
119	SO ₃ ⁻	
120	SO ₃ ⁻	

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-continued

Compound number	R7	R8
121	SO ₃ ⁻	
122	SO ₃ ⁻	
123	SO ₃ ⁻	
124	SO ₃ ⁻	
125	SO ₃ ⁻	

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or

R7 and R8 are selected from the following table:

Compound No.	R7	R8
310	SO ₃ Na	
323	SO ₃ H	
324	SO ₃ H	
325	SO ₃ H	
330	SO ₃ H	
331	SO ₃ H	

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-continued

Compound No.	R7	R8
332	SO ₃ H	
339	SO ₃ H	

2. The β -lactamase inhibitor according to claim 1, wherein R7 and R8 are according to the following table:

Compound number	R7	R8
323	SO ₃ H	
324	SO ₃ H	
325	SO ₃ H	
330	SO ₃ H	

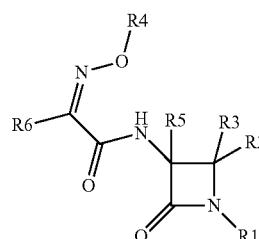
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-continued

Compound number	R7	R8
331	SO ₃ H	
339	SO ₃ H	

3. A pharmaceutical composition, comprising a combination of

a) an antibiologically active compound of the following formula I:



in which

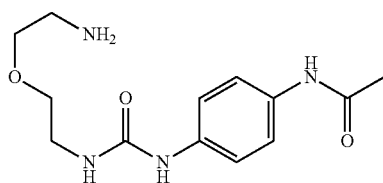
R1 signifies SO₃H, CRaRa'COOH, 5-tetrazolyl, SO₂NHRb or CONHRc,

wherein Ra and Ra' are independently selected from hydrogen; alkyl; allyl; benzyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; phenyl which may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; alkylamino; dialkylamino; alkoxyalkyl and a 5-6 membered heteroaromatic ring which may be substituted with 1 to 4 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

wherein Rb is hydrogen; alkyl; alkoxy; carbonyl; alkylaminocarbonyl; benzylaminocarbonyl in which the benzyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen; or phenylaminocarbonyl in which the phenyl may be substituted with 1 to 5 substituents selected from alkyl, hydroxyl, alkoxy, amino, alkylamino, dialkylamino and halogen;

H) as the compound of formula I of claim 3, the compound (1) wherein R1 is SO₃H, R2 is hydrogen, R3 is CH₃, R4 is (1,5-dihydroxy-4-oxo-1H-pyridin-2-yl)methylene, R5 is hydrogen and R6 is 2-amino-1,3-thiazol-4-yl, and an inhibitor (323) of formula II of claim 1, wherein R7 is SO₃H and R8 is

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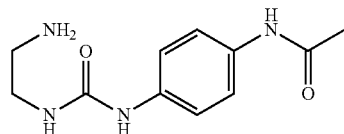
and
sulbactam or clavulanic acid or a pharmaceutically acceptable salt thereof

or

I) as the compound of formula I of claim 3, the compound (1) wherein R1 is SO₃H, R2 is hydrogen, R3 is CH₃, R4 is (1,5-dihydroxy-4-oxo-1H-pyridin-2-yl)methylene, R5 is hydrogen and R6 is 2-amino-1,3-thiazol-4-yl,

and

an inhibitor (324) of formula II of claim 1, wherein R7 is SO₃H and R8 is



and
clavulanic acid or a pharmaceutically acceptable salt thereof

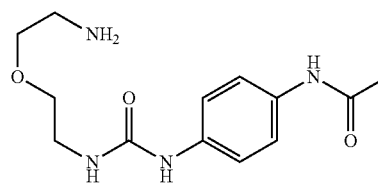
or

O) as the compound of formula I of claim 3, the compound (21) wherein R1 is SO₃H, R2 is hydrogen, R3 is CH₃, R4 is (1,5-dihydroxy-4-oxo-1H-pyridin-2-yl)methylene, R5 is hydrogen and R6 is 4-amino-pyrimidin-2-yl

and

an inhibitor (323) of formula II of claim 1, wherein R7 is SO₃H and R8 is

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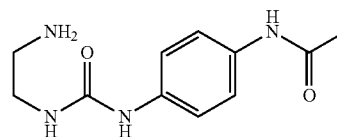
and
sulbactam or clavulanic acid or a pharmaceutically acceptable salt thereof

or

P) as the compound of formula I of claim 3, the compound (21) wherein R1 is SO₃H, R2 is hydrogen, R3 is CH₃, R4 is (1,5-dihydroxy-4-oxo-1H-pyridin-2-yl)methylene, R5 is hydrogen and R6 is 4-amino-pyrimidin-2-yl

and

an inhibitor (324) of formula II of claim 1, wherein R7 is SO₃H and R8 is



and
sulbactam.

5. An article containing a compound of formula I as defined in claim 3, as an agent for treating infections caused by pathogenic Gram-negative bacteria, and an inhibitor of formula II of claim 1, as a combination for the simultaneous, separate or successive administration in the treatment of infections caused by pathogenic Gram-negative bacteria.

* * * * *